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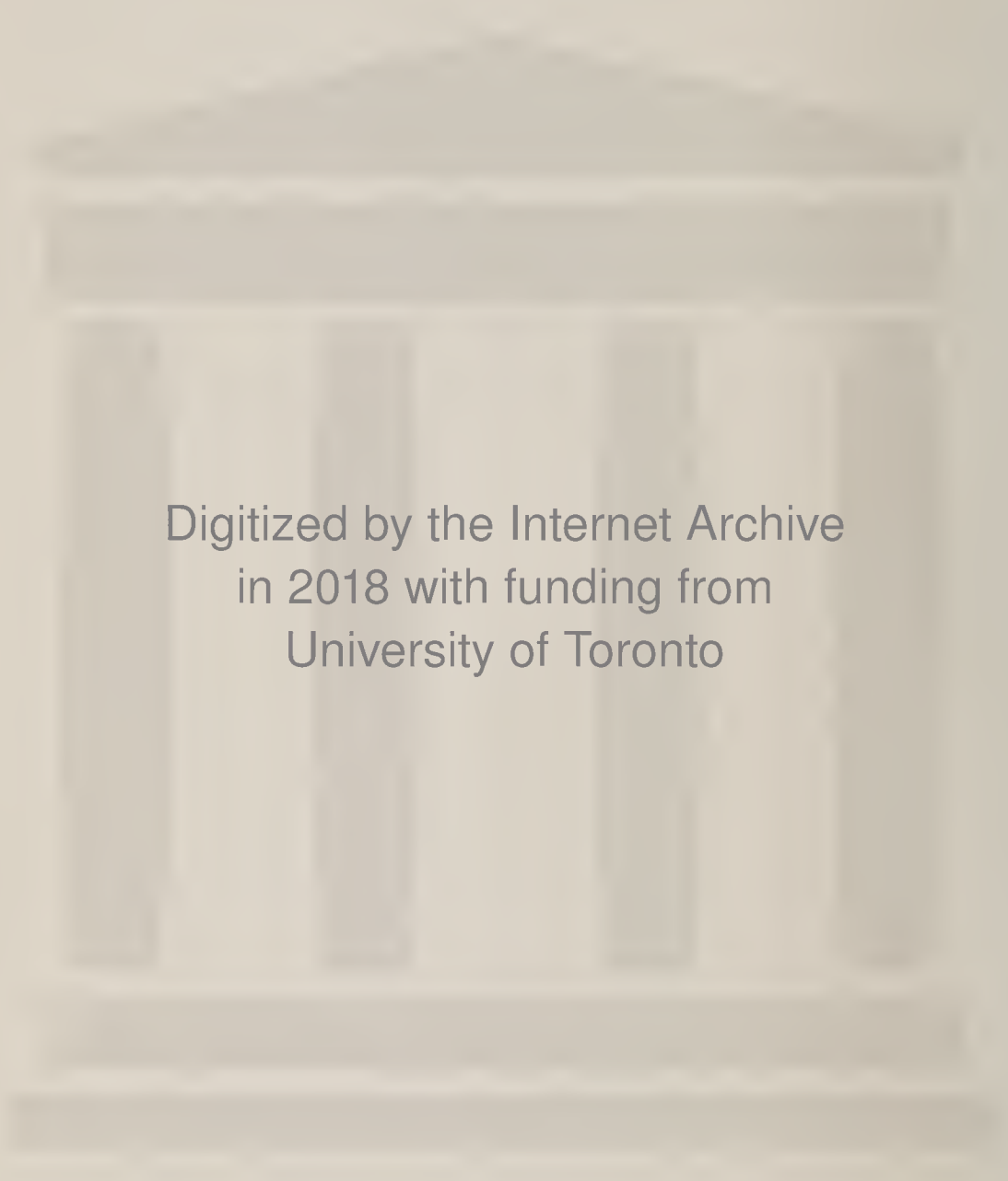












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# HUMAN PHYSIOLOGY





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AN INTRODUCTION  
TO  
HUMAN PHYSIOLOGY

BY  
*desire*  
AUGUSTUS D. WALLER, M.D., F.R.S.  
"

LECTURER ON PHYSIOLOGY AT ST MARY'S HOSPITAL MEDICAL SCHOOL, LONDON

THIRD EDITION

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TO THE MEMORY  
OF  
MY FATHER  
AUGUSTUS WALLER, M.D., F.R.S.

1816—1870

*Emigration of Leucocytes, 1846*

*Degeneration and Regeneration of Nerve, 1850*

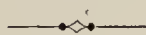
*Cilio-spinal Region, 1851*

*Vaso-constrictor Action of Sympathetic, 1853*





## P R E F A C E



*I have written this book with a distinct consciousness that physiology in a medical school is, in conjunction with anatomy, the introduction to medicine and to surgery of those who will be engaged in 'general practice,' and I hope that the volume may be found to justify its title of 'Human Physiology' with reference to the position that the subject should occupy in medical education, as the junction to which anatomy, chemistry, and physics converge, and from which the principles of medicine and of surgery diverge.*

*Physiology does not consist in a knowledge of recondite phenomena, of difficult names, and of complicated instruments. It should fundamentally consist in the living mental picture of what the great organs below a man's skin are like, what they are doing, how they can be examined, what happens when they are not working properly, how their actions hang together, how they may be influenced for good and for evil. But physiology, as it is written, contains more than this, and, in sparing measure, it is well that the 'Institutes of Medicine' should not be restricted to the visibly 'useful,' or to the obviously 'utilisable.'*

*A short course of practical physiology is of the utmost value to the student of medicine, not merely because it puts into his hands the methods of everyday medical chemistry and physics, but because it helps to correct that credulous bias or primitive 'suggestibility' which is a physiological property of the human brain, and only too apt to be fostered by unmitigated bookwork. Practical physiology is the indispensable adjuvant and corrective of book-physiology,*

because it brings words to the touchstone of physical instruments and of chemical tests, because the conversion of phrases into facts gives measure of their relative value, and is at the same time an exercise of patience and of judgment—in short, because it is a rational discipline, as well as a useful technique.

Nor is this all. An acquaintance with at least the existence of frontier interests, and even a participation *de tactu* with science at its growing surface, cannot be excluded from the laboratories of a medical school without serious and immediate loss to the interests of medicine and of surgery.

With regard to the plan of construction of this book. The chapter headings are intended to serve a double purpose: as a summary or syllabus—being, in fact, the condensed lecture-syllabus that I have used during the last nine years; and as a means of self-examination. A student having read certain paragraphs or chapters, should return to the chapter heading, and find by its recapitulation whether it contains expressions that are meaningless to him, or whether it recalls a series of ideas that he has understood and assimilated. If an unfamiliar technical word is met with, its meaning should be ascertained at some other part of the book, through the index.

Certain subjects are more difficult than others; some are more suitably studied during the first year, others during the second year; others, again, are difficult and of practical importance, some are difficult and of scientific importance; one subject may deserve days or weeks of study and require thorough analytic treatment, or be adapted to experimental demonstration; others may be more lightly passed over. Detailed guidance in such matters will be supplied by the discretion of the teacher, and by the traditions current among examinees; but in order to give some general guidance to the private reader, occasional use has been made of an asterisk (\*) in the chapter headings. In Part I. the sign has been freely used to indicate portions upon which a first-year student should not spend time if he should find the subject difficult. In Part II. the sign has been sparingly used, to indicate portions of undoubted difficulty, that can be omitted by a second-year student who is not reading for the higher examinations. Subjects of unmistakable difficulty in Part



*II.*, such as the brain, vision, &c., have not been so marked, because their great practical importance demands early and patient study from every student of medicine.

The descriptions of figures are to be regarded as forming part of the text, inasmuch as they frequently contain statements of fact that are not included in the text. The Appendix contains matter that was felt to interfere with continuity of description in the main account of physiological phenomena.

I have given a Bibliography after some hesitation, feeling that references to original papers are of no use to junior students, and must be too imperfect to be satisfactory to more advanced students. But to do so has allowed the text to be lightened of names, while preserving for the student who may inquire further, the authorities for the principal statements made. Moreover, the chief use of a bibliography in a general text-book is to afford a few main starting-points and indications, to be followed up in the year-books and archives of a well-equipped library; and even for junior students, it ought to be of some assistance to know whether doctrines and statements date from last year or from last century. The year of publication has been given in each case, leaving the particular page of a particular paper to be looked up in the index of the journal to which reference has been made.

The preparation of new editions has given me the opportunity of making several corrections; I have also made some alterations and transpositions of text that seemed to me to be desirable. Attention has been paid to recent work, but I have felt that the gradually-formed deposit of accepted knowledge must be of greater intrinsic value than the latest 'discovery' or the newest theory. An early mental diet in which these items are predominant is an unwholesome diet; their function in elementary instruction is that of condiments, valuable only in conjunction with a foundation of solid food, serving above all to teach that the 'Institutes of Medicine' are not a museum of dead dicta, but a living knowledge, of which the verbal formulæ are unstable—least unstable in the older strata, most unstable at the growing surface. I have therefore deliberately abstained from mentioning many matters that I have heard of or

*read of, and such recent investigations as are mentioned, I have for the most part either worked at myself or seen worked at by others. The principal changes in this edition are in Chapters X and XI; these, as well as other statements for which I am directly responsible, as on pp. 374, 394, 401, are indicated by the letter (w.).*

A. D. WALLER.

WESTON LODGE, 16 GROVE END ROAD, LONDON, N.W.

*September, 1896.*



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# HUMAN PHYSIOLOGY

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## PART I

### *THE PHENOMENA OF NUTRITION*

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#### CHAPTER I

##### *INTRODUCTION*

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- 1 The scope of physiology—Living matter—Metabolism—Up and down—Food and energy—Supply—Expenditure—Death—Organisation—Division of labour.
- 4 The tripod of physiology: anatomy, chemistry, and physics—Its methods of study—Amœba to man—The composition of a man's food—The composition of a man's body—The composition of protoplasm—Six proximate principles—Three important elements—The proteid molecule—Proteid and protoplasm—Comparison of protoplasm, body-stuff, food-stuff, milk, and blood.
- 8 Oxygen and carbon dioxide.
- 8 Excitability.

PHYSIOLOGY deals with the chemical and physical changes that occur in and by living matter. Vegetable physiology deals with plants, animal physiology with animals, human physiology with man.

We cannot define 'life' in physical terms; we can, however, observe and state in what essential particulars living differs from non-living matter, or from matter that is dead.

The essential feature of living matter is its instability; it is the seat of chemical changes, collectively termed metabolism.<sup>1</sup> These changes are divisible into—1. Constructive, integrative, anabolic, or synthetic processes, in the course of which non-living matter is annexed or assimilated by living matter; 2. Destructive,

<sup>1</sup> μεταβολή, change; ἀνά, up; κατά down.

disintegrative, katabolic, or analytic processes, in the course of which living matter and storage substances are expended.

Living matter is supplied by food, the potential energy of which is utilised in the course of analytic changes. These changes mainly fall upon storage substances, and only involve living substance to a very small extent; otherwise expressed, the greater portion of the food is consumed under the influence of living substance without ever forming an integral part thereof.

At any given moment a living body, considered simply as a mass of living matter, possesses a fund of potential energy derived from the previous storage and assimilation of foreign matter; this potential or stored-up energy is constantly flowing off in the form of actual energy as heat and as mechanical work; it is constantly replenished by the continued assimilation of foreign matter. Sooner or later restoration ceases, energy runs down, and the living body ceases to live.

But a living body is not merely a homogeneous mass of living matter; the living matter or protoplasm has become differentiated and organised, and it has surrounded itself with protoplasm products. The lowest form of living matter is a simple cell, *e.g.* an amœba or a leucocyte. The highest form of living matter is the human body, which is literally a 'nation' of cells derived from a single cell called the ovum, living together, but dividing the work, transformed variously into tissues and organs, and variously surrounded by protoplasm products.

In all living organisms a *physiological division of labour* is associated with *anatomical differentiation of structure*. An organism ranks high or low in the animal scale in accordance with the greater or less diversity of its tissues and functions; and in every organism above the lowest, portions of various tissues are found united into distinct masses forming *organs* and possessing *functions*—each organ served by and serving the entire organism. The organism is a community; its individuals are cells; groups of its individuals are organs. In any animal above the lowest we distinguish from a physiological point of view the following principal tissues:—

1. Blood,
2. Epithelium,
3. Connective tissue,
4. Nervous tissue,
5. Muscular tissue,

and attribute them to the following principal functions:—



The digestive preparation and absorption of food-material, and the rejection of waste matter *by epithelium*, the cells of which form the essential parts of secretory glands, of absorbent surfaces, and of excretory glands.

The distribution of food-material and removal of waste material *by blood*.

Internal and external mechanical movements *by muscle*.

Control of action, and communication between parts *by the nervous tissues*.

Mechanical support, connection and preservation of form *by the connective tissues*.

In the vertebrate type of animal we distinguish from a physiological point of view the following principal organs:—

1. of digestion,
2. of respiration,
3. of circulation,
4. of excretion,
5. of reproduction ;

and we collectively consider as ‘organs’—

6. The muscular system, or organ of movement.
7. The nervous system, or organ of control.

The *organ of digestion* comprises the alimentary canal and annexed secretory glands, and has as its function the preparation and absorption of food. Its essential and active element is the epithelial cell.

The *organ of respiration* is the lung, and has as its function the exchange of gases between the organism and the atmosphere.

The *organ of circulation* comprises the heart and vessels with their fluid contents ; its function is the distribution and renovation of blood throughout the organism. Its essential element is muscular fibre.

The main *organ of excretion* is the kidney, which, however, is supplemented by the action of the lung, the skin, and the alimentary canal ; its principal function is the separation of *urea* from the blood ; its essential and active element is the epithelial cell.

The *organs of reproduction* are the ovary and the testis ; their essential elements are the ovum and the spermatozoon respectively, both of which are originally derived from epithelial cells.

The *muscular and nervous systems* conjointly form an organ which is at the same time master and servant of the entire organism. The government of the body is of the constitutional

type; nerve-centres acted upon through nerves receive information of the external conditions and of the internal wants of the entire organism, and, acting through nerves, bring about the movements that satisfy the demands made. The principal nerve-centres are the brain and the spinal cord. The principal sense organs are the skin, the eye, the ear, the nose, and the tongue. The muscles, or executive organs, are distinguishable into muscles of external movements and muscles of internal movements; the former act upon bones, and effect movements directed to or from external circumstances, the latter surround cavities, and effect the internal movements requisite to the transport of matter within the organism. We shall return to the nervous and muscular systems in Part II., and shall then consider their functions at length.

Human physiology, or the knowledge of the functions of the cells, tissues, and organs that constitute the body, requires, in the first place, an elementary knowledge of anatomy, of chemistry, and of physics. By anatomy, including microscopical anatomy, we learn what the cells, tissues, and organs are like, where they are situated, what channels lead to and from them. By chemical methods we learn what they are composed of, what they make, and what their products can do. By physical methods we learn what work they do in and out of the body, and how their functions are modified by external forces.

Given these three essentials, physiological processes and consequences are studied by exact observation and by insulative experiment, upon the lower animals, upon the higher animals, and upon man. The lower animals, the frog in particular, are especially suitable to the study by direct experiment of the functions of isolated tissues, such as nerve and muscle. The higher animals—rabbit, cat, dog, horse, monkey, &c.—are required for the study of organs which are anatomically similar to those of man. On man we carry out such experiments as require intelligent co-operation on the part of the subject, and we record exact observations concerning the natural history of disordered functions, and the effects upon such functions of various remedial interferences.

The complicated functions of the human body are the highest expression of physiological activity, the functions of the simple protoplasmic cell are its lowest and earliest manifestations. The following general facts apply to all living matter; they are observable in the everyday life of a man; they are logically



attributable to the smallest lump of protoplasm, such as a single white blood-corpuscle.

The following articles—*meat* (with *fat*), *bread*, *salt*, *water*, and the *air* we breathe—contain representatives of all the various classes of proximate principles which must enter a man's body to keep that body living and energy-yielding. The meat contains the proximate principles<sup>1</sup> *proteid* and *fat*. The bread contains *carbohydrate*; to these are added *salt* and *water*, and from the air *oxygen*. These six proximate principles constitute the essential income of all living matter—of each microscopic speck of protoplasm as of the entire animal body. The three most important elements which they contain are *oxygen*, *nitrogen*, and *carbon*.

Oxygen we need not at present consider beyond remarking that it is a simple element, a gas, which easily finds its way into and is assimilated by living matter.

*Carbohydrates*, *fats*, and *proteids* are the three groups that most demand our attention at this stage. They all contain the elements, carbon, oxygen, and hydrogen; proteids contain in addition the element nitrogen. The entire nitrogen income is contained in proteids; the chief carbon income is supplied in carbohydrates and fats. A glance at their formulæ shows further some noteworthy points.

*Carbohydrate*.—The formula of the carbohydrate, dextrose, is  $C_6H_{12}O_6$ . All carbohydrates, in common with dextrose, contain no nitrogen, and they contain hydrogen and oxygen in proportion by volume to form water, viz.  $\frac{2 \text{ hydrogen}}{1 \text{ oxygen}}$ . If a carbohydrate is burned, *i.e.* completely oxidised, it requires additional oxygen from outside itself only for the carbon; the amount of oxygen in the carbohydrate itself is sufficient for the complete oxidation of the hydrogen it contains.

*Fat*.—The formula of the fat, tripalmitin, is  $C_{51}H_{98}O_6$ , or  $C_3H_5 \cdot 3(C_{16}H_{31}O_2)$ . Fats contain no nitrogen; they contain relatively to the proportion  $\frac{2 \text{ hydrogen}}{1 \text{ oxygen}}$ , a great deal of hydrogen, very little oxygen. In tripalmitin, for instance, the proportion of hydrogen to oxygen is much higher,  $\frac{16 \text{ hydrogen}}{1 \text{ oxygen}}$ . To be completely oxidised, fat requires outside oxygen to satisfy hydrogen as well as carbon.

*Proteid*.—The formula of a proteid is given as  $C_{72}H_{112}N_{18}O_{22}S$ , but this is not a true chemical formula, proteids, however pure,

<sup>1</sup> A proximate principle is any element or group of elements that can be separated as such without decomposition from animal or vegetable bodies.

being of variable composition. We see, however, by this approximate formula that a proteid contains nitrogen, and that the proportion between hydrogen and oxygen is greater than in starch, less than in fat, viz.  $\frac{5 \text{ hydrogen}}{1 \text{ oxygen}}$ . By weight the percentage composition of various proteids is

$$\text{C.} \begin{cases} 51.5 \\ \text{to} \\ 54.5 \end{cases} \quad \text{H.} \begin{cases} 6.9 \\ \text{to} \\ 7.3 \end{cases} \quad \text{N.} \begin{cases} 15.2 \\ \text{to} \\ 17 \end{cases} \quad \text{O.} \begin{cases} 20.9 \\ \text{to} \\ 23.5 \end{cases}$$

Moreover, the proteid molecule holds to itself, if not within itself, small, but important, quantities of *sulphur*, of *phosphorus*, and of *iron*. There is sulphur in every proteid, however pure; there is phosphorus in most proteids, and especially in that of cell-nuclei; there is iron in one most important compound, hæmoglobin.

A proteid molecule is very large, and composed of a large number of atoms of C. O. H. N.; these may 'hold hands,' so to speak, in many different ways; the molecule, while forming part of protoplasm, may take many shapes, may offer many kinds of opportunities for different kinds of chemical intercourse. It is to this character that the proteid molecule owes its preponderating importance in living matter; the molecule, while in living matter, is 'all alive,' the seat and centre of an active commerce of atoms.

**The composition of protoplasm.**—The relation between proteid and protoplasm is very close; proteid, as we examine it chemically, is non-living, protoplasm is living; if we attempt to examine protoplasm chemically, we necessarily kill it in the attempt. But if, having killed protoplasm, we continue the chemical examination, we find that the killed matter is mostly proteid and water. It appears that water is an integral part of protoplasm. Pushing the examination still further, we find that our killed protoplasm contains a small proportion of carbohydrate, a small proportion of fat, and a small proportion of salts.

Thus we have found, in the analysis of protoplasm, that its main bulk is composed of water and proteid, that it also comprises some carbohydrate, some fat, and some salt. And by this different channel we are brought to our starting point. The list just given is the same as that already given of the proximate principles required to keep a man alive, with the one exception of oxygen, which we did not find in our analysis of dead proto-



plasm. This exception is, however, easily accounted for; we shall find that a supply of oxygen is necessary to protoplasm, but that we need not expect to find any free oxygen left in killed protoplasm. We may sum up these considerations in the following form :—

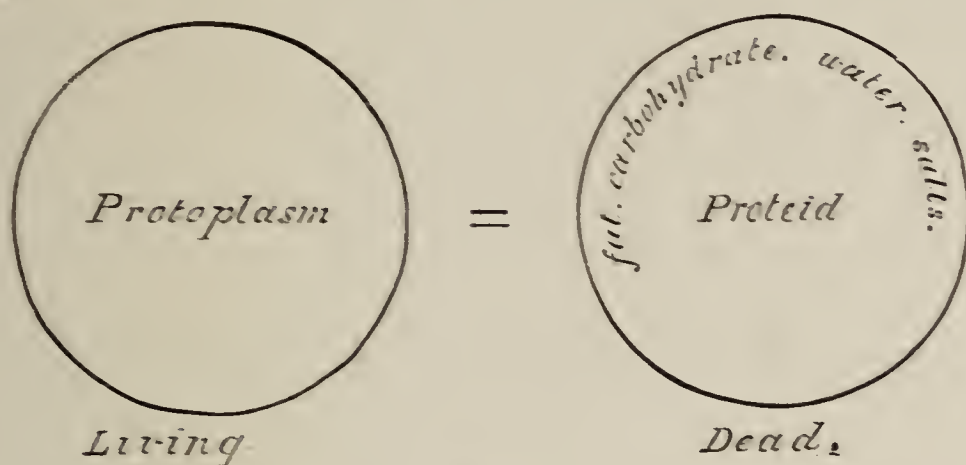


FIG. 1.

The chemical elements contained in the proximate principles above considered are C. O. H. N. They enter the body in *proteids*, *fats*, and *carbohydrates*, and together with *water*, *salts*, and free *oxygen* they concur to the formation and maintenance of *protoplasm*.

The composition of the body.—Knowing what goes into the body, we have a general knowledge of its composition; its chief component elements will be C. O. H. N.; its chief component proximate principles will be carbohydrates, fats and proteids, water and salts.

*Water* forms at least two-thirds of the total body-weight. A human body weighing twelve stone if completely dried would weigh about four stone. After cremation the residual ash is 2 to 3 per cent., *i.e.* three to five pounds. The proportion of water varies in the various tissues and fluids; it is greatest in the saliva and in gastric juice, least in the enamel of the teeth, but even in bones the proportion is considerable. The following are representative numbers :—

	Water per 1,000	Solids per 1,000
Saliva . . . . .	995	5
Gastric juice . . . . .	973	27
Milk . . . . .	891	109
Blood . . . . .	791	209
Muscle . . . . .	757	243
Bone . . . . .	486	514
Enamel of teeth . . . . .	2	998

*Salts* are contained in all the tissues and fluids. The principal salts are :—Sodium chloride, in the fluids especially; potassium

phosphate, in the tissues especially ; calcium phosphate, in bone particularly. The above (viz. water and salts) are the chief inorganic compounds ; the organic compounds are *proteids*, *fats*, and *carbohydrates*, which we have considered sufficiently at the present juncture, but shall refer to again hereafter.

We may now realise the essential similarity between the composition of protoplasm and that of the entire human body, and of its normal food-supply. Each comprises the same elements and the same proximate principles ; we shall find that *milk*, which is the sole food in early life, and *blood*, which is the general food of the whole body—and during gestation the fund from which a new body is entirely formed in the mother's womb—likewise comprise the six proximate principles enumerated above.

**Oxygen.**—We have left for separate consideration this necessary item of supply. A man must breathe to keep alive ; protoplasm runs down and dies if deprived of oxygen. The tissues of the human body get oxygen from the blood, which gets it by the lungs. The blood is oxygenated at the lungs, deoxygenated or reduced by the tissues. Protoplasm gets oxygen from the air, or from oxygen-holding compounds which it can decompose. This deoxygenating or reducing action of protoplasm is a fundamental characteristic of the living state. Living matter requires oxygen, and gets it from the air, or by taking it away from combinations which do not hold it fast. And although oxygen is continuously necessary to living matter, we do not at any given moment find free oxygen in protoplasm or in tissue. It is at once drawn into chemical combination. It reappears afterwards in company with carbon, as *carbon dioxide*, which must be discharged. A man exhales carbon dioxide, protoplasm exhales carbon dioxide. But, as we shall see hereafter, the carbon dioxide comes, not from any direct union of free oxygen with free carbon, but indirectly from complex compounds in which the incoming oxygen had become stored.

The signs by which we can always recognise that protoplasm is alive are—(1) its reducing or deoxygenating power, (2) its exhalation of carbon dioxide, (3) its **excitability**. The first two tokens have just been considered. The third—**excitability**—manifests itself as movement in response to excitation, and will claim our attention more particularly when we come to the study of the nervous system. At this stage it will be sufficient for us to recognise 'excitability' as a fundamental property of living matter.



## CHAPTER II

## THE BLOOD

## PAGE

- 9 **Function and composition:** Proximate principles—Elements.
- 10 **Physical properties:** Consistence—Colour—Opacity—Specific gravity—Reaction—Taste—Smell—Coagulability.
- 12 **Microscopic structure:** Corpuscles, red and white—Leucocytes—Hæmatoblasts—Numeration of corpuscles—Hæmacytometer.
- 13 **Chemical composition.**
- 15 **Plasma and serum: fibrinogen and fibrin:** Preparation of serum, of fibrin, of defibrinated blood, of plasma—\*Centrifugal apparatus—Magnesium sulphate plasma—Cold plasma—‘Peptone’ plasma—Buffy coat—Lake blood.
- 17 **The composition of plasma, lymph, chyle, and serous fluids.**
- 18 **Facts and theories relating to coagulation:** Hydrocele fluid and serum—Buchanan—Schmidt—Hammarsten—\*Present state of the question.
- 21 **Methods of separating the plasma proteids:** Fibrinogen—Serum-globulin—Fibrin-ferment—Serum-albumin.
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- 24 **The life-history of the blood-corpuscles:** The red corpuscles: origin, function, end—The white corpuscles: origin, vital characters, functions—Emigration—Suppuration—Cicatrization—Phagocytosis.
- 27 **The blood-gases:** \*The mercurial gas-pump.
- 29 **Hæmoglobin—its compounds and derivatives:** Normal, abnormal, and artificial—The spectroscope.
- 35 **Arterial blood and venous bloods:** Hepatic blood—Portal blood—Blood from glands and from muscles—Temperature; cooled venous blood—warmed venous blood.
- 37 **Alterations of the blood in disease:** Anæmia—Leucocythemia—Inflammatory blood—Kidney affection—Gout—Liver affection—Diabetes—Hæmophilia.
- 38 **Quantity of blood in the body:** Its estimation.
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**Function and composition.**—A regular supply of healthy blood is an essential condition of the life of the animal body, and of its several parts. The whole body dies if blood be lost in excessive quantity; a single organ, or a single limb, dies if its supply of blood be permanently arrested. The blood is the medium of nourishment to the tissues of the body, and contains, therefore, all the materials that they require, viz. *oxygen, water,*

*proteids, carbohydrates, fat, salts.* Living tissues are constantly in a state of chemical change, consuming and assimilating to themselves materials which they derive from the blood, and rejecting the consequent waste products. These are discharged into and carried away by the blood, which contains, therefore, the materials in their used-up form, viz. *carbonic acid* and *urea*.

In correspondence with the large proportion of proteids and of hæmoglobin contained in blood, the elementary composition of its solid residue agrees closely with that of proteids in general, and with that of muscle, which is essentially proteid. A comparison of some actual analyses, giving the amounts of the chief elements entering into the composition of blood, muscle, hæmoglobin, and proteids, will best illustrate this similarity:—

Elementary composition (in the dried state)	Of Blood	Of Muscle	Of Hæmoglobin	Of Proteids
Carbon . . . .	51.96	51.86	53.8	51.5 to 54.5
Oxygen . . . .	21.30	21.30	21.2	20.9 to 23.5
Hydrogen . . . .	7.25	7.58	7.1	6.9 to 7.3
Nitrogen . . . .	15.07	15.03	16.1	15.2 to 17

**Physical properties.**—Blood in the body, or freshly drawn, is a slightly viscid fluid, bright scarlet in colour if taken from an artery, dark purple or almost black if taken from a vein; in either case opaque, even in thin layers. It is heavier than water, its *specific gravity* being about 1,050 as compared with water, 1,000—*i.e.* 1,000 cubic centimeters of water weigh 1,000 grammes, 1,000 cubic centimeters of blood will weigh about 1,050 grammes;<sup>1</sup> its *reaction* is alkaline, never acid—the popular expression ‘acidity of the blood’ has no foundation in fact. The alkalinity varies in degree; it is at a maximum after meals, owing to the secretion of acid gastric juice, and at a minimum after severe exertion, owing to the acid which is formed in muscular contraction; the alkalinity of blood is attributable to the presence

<sup>1</sup> Measurements will be given throughout according to the metric system.

1 gramme = about 15 grains.

100 cubic centimeters, or 100 c.c. = about  $3\frac{1}{2}$  fluid ounces.

For microscopic measurements the unit is the micromillimeter, or  $1\mu = \frac{1}{1000}$  millimeter = about  $\frac{1}{25000}$  inch.

The preliminary difficulties generally experienced by students in the use of metric measurements are soon overcome, and their greater simplicity appreciated. The relative magnitudes  $8\mu$ ,  $10\mu$ ,  $15\mu$ ,  $50\mu$  are far more readily realised than the corresponding values  $\frac{1}{3200}$  in.,  $\frac{1}{2540}$  in.,  $\frac{1}{1600}$  in.,  $\frac{1}{500}$  in.

The capacities of vessels used in the practical classes are marked on the metric system, and students soon learn that a 100 c.c. flask holds rather less than a 4-oz. bottle, and that a 500 c.c. beaker is the equivalent of a short pint.



of disodic phosphate— $\text{Na}_2\text{HPO}_4$ , and the degree of alkalinity is about the same as that of a .3 per 100 solution of sodium carbonate, *i.e.* equal to a  $\frac{\text{normal}}{20}$  alkaline solution. In consequence of the presence of sodium chloride, blood has a saltish taste, and it possesses a faint and indescribable smell, which is made more distinct by the addition of a few drops of sulphuric acid. The property that is the prominent characteristic of blood is its *coagulability*. Left undisturbed for a few minutes after it has been drawn, the fluid blood becomes more and more viscid; in a few minutes it is no longer a fluid, but a red jelly; as time goes on, the jelly shrinks and gets firmer, and squeezes out drops of almost colourless liquid; in a few hours the jelly has shrunk to

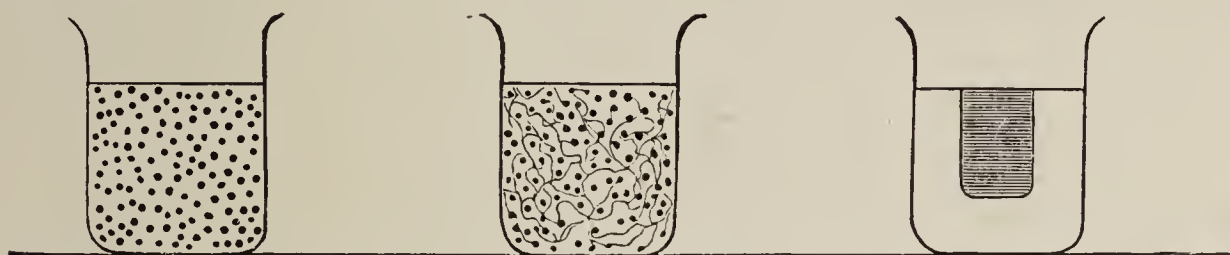


FIG. 2.—DIAGRAM TO ILLUSTRATE THE PROCESS OF COAGULATION.

I. Fresh.

II. Coagulating.

III. Coagulated.

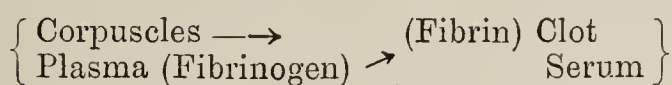
(Corpuscles and plasma.)

(Birth of fibrin.)

(Clot and serum.)

Plasma minus fibrinogen equals serum. Corpuscles plus fibrin equal clot.

its utmost, and floats in a considerable quantity of the liquid. Coagulation has taken place and is now complete, the shrunken jelly is the *clot*, the liquid in which it floats is the *serum*; coagulated blood consists of clot and serum. The living blood from which this clot and serum were formed consisted originally of corpuscles and plasma, and, as we shall presently learn, the change is essentially a change of the plasma, fibrin arising in it from fibrinogen, gathering itself together and sweeping the corpuscles from the fluid.



Thus the plasma minus fibrinogen becomes serum, and fibrin plus the corpuscles forms the blood clot.

The process of coagulation is hastened by moderate warmth, and by contact with foreign bodies; it is retarded by cold, by certain salts (sodium and magnesium sulphate in particular), and when the blood is protected from contact with foreign bodies by means of oil (with which it does not come into actual contact). Blood that has been frozen before coagulation has taken place may be so kept for months, and yet will clot when thawed; this



fact implies that coagulation is not a vital act, but only a physico-chemical change.

**Microscopic structure.**—A small drop of blood obtained by pricking the finger, and quickly covered for examination under the microscope, is seen (with a magnifying power of 300 diameters) to be composed of a multitude of corpuscles floating and rolling over and over in the minute eddies of a fluid which is not itself visible—the *liquor sanguinis* or *plasma*. Examined more closely, the corpuscles will be seen to be of two kinds—the *coloured corpuscles*, the more numerous, and the *white corpuscles*, far fewer in number, 2 or 3 of the latter among 1,000 of the former, so that they require to be looked for with some little care. All the more so that the tint of the coloured corpuscles is faint, they are not far from colourless, and it is only by their great number that the strong red colour of blood is produced. On still more careful examination with very high powers, very small colourless corpuscles—about  $2\mu$  in diameter, 2 or 3 to every 100 red corpuscles—may be distinguished; these are called

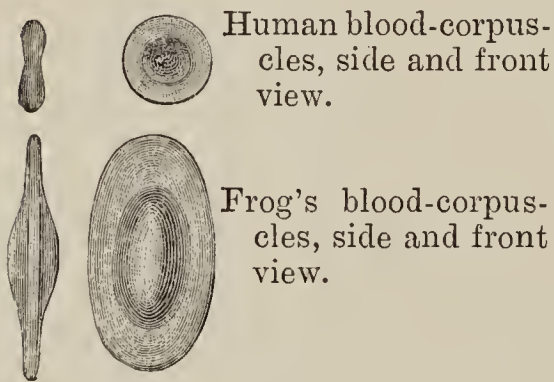


FIG. 3.

The corpuscles are drawn to the same scale, viz.  $\times 1000$ .  $1\mu$  or  $\frac{1}{1000}$  mm. of actual length is represented by 1 mm.; the human corpuscle as sketched above is  $7\mu$  in diameter, i.e. its actual diameter is  $7\mu$ ; the frog's corpuscle is  $20\mu$  by  $11\mu$ . The corpuscles of mammalia resemble those of man, but differ in size. Average magnitudes are: elephant,  $9\mu$ ; dog, rat,  $7\mu$ ; cat, horse, ox,  $6\mu$ ; goat, sheep,  $5\mu$ . But the corpuscles of man, averaging  $7\mu$ , may range from as low as  $5\mu$  to as high as  $9\mu$ .

*blood-platelets* or *hæmatoblasts*. A single coloured corpuscle seen flat is circular, and, as the microscope is focussed up and down with the fine adjustment, appears now as a dark centre surrounded by a light ring, now as a light centre surrounded by a dark ring. This appearance is not due to a nucleus—human blood-corpuscles, in common with those of all mammalia, have no nucleus—but to the cupped shape of the corpuscles; a single corpuscle seen sideways looks something like a dumb-bell, and the two views combined show that the actual shape of the corpuscle is that of a disc cupped on both surfaces—a bi-concave disc; these discs frequently run together and adhere to each

other in longer or shorter rolls like piles of coins; if distilled water be added to blood, water enters the red corpuscles and causes them to become globular and then burst; if strong solution of salt be added, water leaves the corpuscles, which become shrivelled

and crenated ; these experiments indicate that the corpuscle is surrounded by a film. The blood of amphibia, reptiles, fishes, and birds has corpuscles different from those of mammalian blood in shape and structure. Of all these lower classes the red corpuscles are oval, convex on both sides, and their central part differs in structure from the remainder of the corpuscle, constituting a nucleus.

Various instruments have been devised for enabling an estimate to be formed of the number of corpuscles present in a given volume of blood ; the principle of all such instruments is to furnish the observer with a very minute cubic space under the microscope, in which it is possible to see and count the corpuscles of blood diluted a given number of times. The form of instrument most used in this country has been called a *hæmacytometer* (Gowers) ; the blood is diluted 200 times, and observed through the microscope in a cell  $\frac{1}{5}$  millimeter deep, and divided into  $\frac{1}{10}$  millimeter squares : the cubic space of each division is thus  $\frac{1}{500}$  millimeter, and the number of corpuscles in one division multiplied by 100,000 gives, therefore, the number of corpuscles per cubic millimeter. Normally, this number is 4,000,000 to 5,000,000 (40 to 50 in each square) ; abnormally, in extreme anæmia, it may be as low as 1,000,000 to 2,000,000 (10 to 20 in each square).

The white corpuscles, or leucocytes, are of all shapes while they are alive, and their shapes are constantly, but very slowly, changing ; ultimately—*i.e.* as they die—they assume a globular form. Many of them break up and vanish. They are of various sizes ; some are no larger than the coloured corpuscles, some are even smaller, some are considerably larger ; and of the larger leucocytes, some contain a coarsely granular substance, staining deeply with acid dyes, such as eosin, others are hyaline with a single large nucleus, others still—and these are the most important and numerous—have an irregularly lobed nucleus, staining deeply with basic dyes, such as methyl blue or fuchsin. These last may be regarded as typical leucocytes, and exhibit the various life-characters that will presently be described (p. 25).

**Chemical composition.**—100 grammes of blood consist of about 80 grammes of water and 20 grammes of solids, some in solution, some in suspension, and contain 60 cubic centimeters of gases, some in solution, some in loose chemical combination with other constituents of the blood.



The twenty grammes of solids comprise :—

Hæmoglobin	. . . . .	10 grammes
Proteids	. . . . .	9 „
Fat	}	1 gramme
Carbohydrate		
Salts		
Urea		

The important proteid, *fibrinogen*, forms only a very small proportion of the total proteid ; although the fibrin which it forms appears of considerable bulk when it is separated by whipping, yet its actual weight is small—1,000 grammes of blood yield between 2 and 4 grammes of fibrin. The *hæmoglobin* is contained in the coloured corpuscles, which owe to this substance their function as oxygen-carriers to the tissues. The proteids comprise *serum-albumin* and *serum-globulin* in about equal proportions dissolved in the plasma.

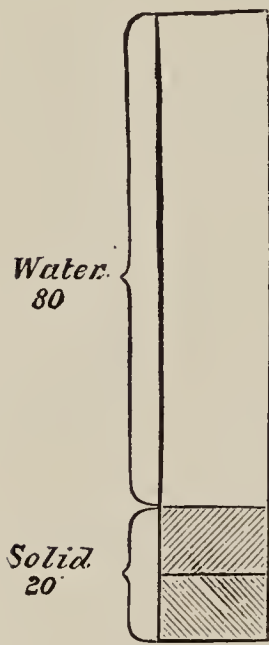


FIG. 4.—THE PRO-  
PORTION OF SOLIDS  
TO WATER IN THE  
BLOOD GRAPHI-  
CALLY EXPRESSED.

Physically considered, 100 grammes of horse's blood consist of about :—

Corpuscles	. . . . .	33 grammes
Plasma	. . . . .	66 „

The 33 grammes of corpuscles comprise : —

Water	. . . . .	22 grammes
Solids (chiefly hæmoglobin)	. . . . .	11 „

The average amount of hæmoglobin in human blood is somewhat higher—between 12 and 15 per 100.

The 66 grammes of plasma comprise :—

Water	. . . . .	60 grammes
Solids	{ Serum-albumin } { Serum-globulin }	. . . . . 6 „

The chief *salts* of the blood are chlorides and phosphates, combined with sodium and potassium. Sodium chloride preponderates in the plasma ; potassium phosphate preponderates in the corpuscles. Other salts present are sulphates and sodium bicarbonate (NaHCO<sub>3</sub>).

*Fat*, *sugar*, and *urea* are constantly present in the blood, but in small quantities only. Fat is present in variable quantity ; after a meal of fat it may be so abundant as to form a scum on the surface of drawn blood ; it remains, however, but a very short time in circulating blood, being separated by the tissues and stored in them ; 1,000 grammes of blood contain from 2 to 6

grammes of fat. Sugar is present in small but constant quantity; 1,000 grammes of blood contain about 1 gramme of sugar. Urea is present in small quantity; 1,000 grammes of blood contain about  $\frac{1}{4}$  to  $\frac{1}{2}$  gramme.

**Plasma, serum, fibrinogen, fibrin.**—The fluid in which the corpuscles are suspended while the blood is uncoagulated is called the *plasma*, or *liquor sanguinis*. It differs from the fluid in which the clot floats after the blood has coagulated, *i.e.* the serum, in this important respect, that plasma contains a body called *fibrinogen*, while serum contains no fibrinogen; both fluids have a specific gravity of about 1,030, and contain about the same quantity of proteids. Fibrinogen is, as its name implies, a body that becomes or gives rise to fibrin, and it is the conversion of fibrinogen into fibrin that is the essential event in the coagulation of the blood. Fibrinogen in the plasma becomes fibrin,

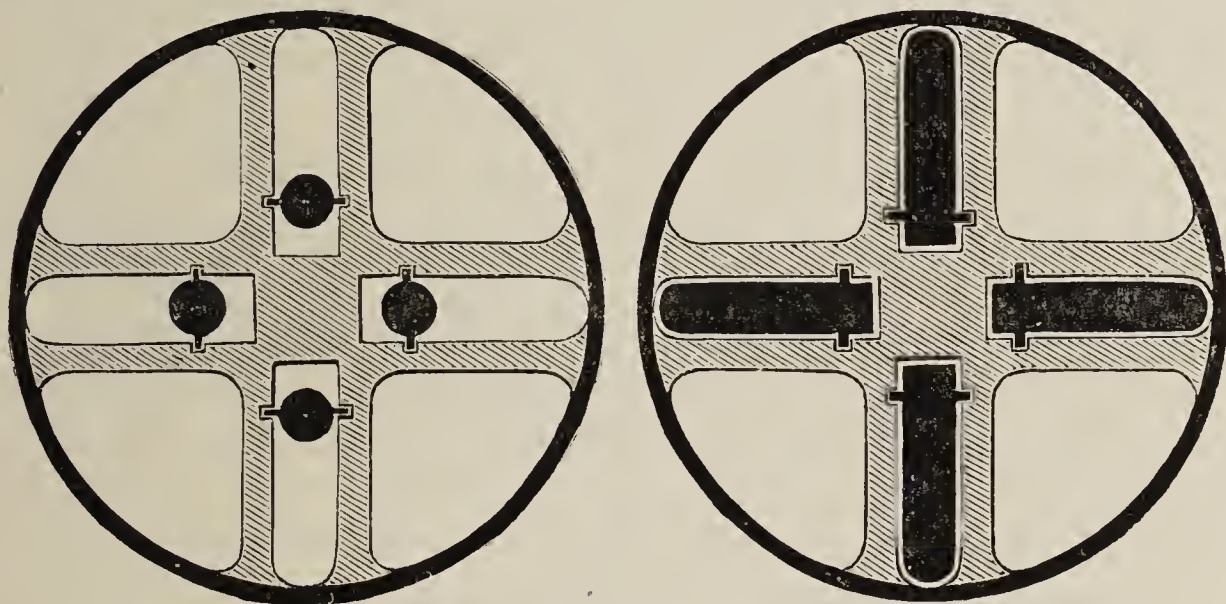


FIG. 5.—PLAN OF CENTRIFUGE AS USED FOR THE SEPARATION OF PLASMA.

**Centrifugal Apparatus.** Disc fitted with four long buckets pivoted so that they hang vertically when the disc is at rest, and assume a horizontal position as the disc revolves. Large test-tubes containing the blood-mixture fit into the buckets. By centrifugal force, while the disc is revolving, the corpuscles collect towards the circumference, *i.e.* at the bottom of the test-tubes: the plasma is left nearer the centre, *i.e.* at the top of the test-tubes.

**Stationary.** The buckets (represented in transverse section) hang vertically from the disc.

**Revolving.** The buckets (represented in longitudinal section) swing out to a horizontal plane.

which pervades the fluid in all directions in the form of innumerable filaments; the plasma having lost fibrinogen, serum is left. The filaments of fibrin shrink, and, entangling in their meshes the corpuscles, sweep these from the serum, and form with them the clot, which gradually shrinks and squeezes out the serum. To obtain *serum* it is only necessary to allow freshly drawn blood to coagulate without disturbance.



To obtain *fibrin*, blood is whipped with a bundle of twigs as soon as it is drawn ; the fibrin adheres to the twigs and forms a stringy mass, which is to be subsequently washed in water and left under a running tap until almost white. The whipped blood from which fibrin has been separated is *defibrinated blood*.

To obtain *plasma* it is necessary to prevent coagulation, so that the corpuscles may subside and leave an upper layer of uncoagulated plasma. To this end, the blood is allowed to flow into a vessel surrounded with ice, and containing a saturated solution of magnesium sulphate. After a few hours the corpuscles will subside if the vessel is left undisturbed, or if, instead of leaving the separation to take place by the action of gravity, the vessel is fixed to a horizontal wheel revolving at a high speed, the separation will be much accelerated by centrifugal force.

It is convenient, but not absolutely necessary, to employ both ice *and* magnesium sulphate. From blood mixed at once with magnesium sulphate at ordinary temperature, plasma can be obtained by subsidence, or by the centrifuge. By cooling blood sufficiently rapidly, and leaving it to stand surrounded by melting ice, plasma may be obtained without magnesium sulphate. Horse's blood, which coagulates more slowly than that of other mammals, is the best kind of blood to choose for the purpose.

Another convenient method of obtaining plasma is afforded by the property which commercial peptones possess of preventing coagulation when they are injected into the vessels (pp. 23 and 189). The blood of a dog so treated does not coagulate after it is drawn, and plasma may be obtained from it by the centrifuge.

The upper part of the clot that forms in normal horse's blood is not red like the lower portions, but pale or buff-coloured. This difference is due to the fact that the red corpuscles of horse's blood cling together in little clumps which quickly subside from the upper part of the fluid, and the clot that forms subsequently is comparatively free from them. A buffy coat has also been noticed in human blood in inflammatory conditions ; the coagulation of normal human blood is too rapid to allow of the formation of a buffy coat, but inflammatory blood coagulates more slowly, and the upper part of the clot is left pale.

Defibrinated blood is opaque, and viewed by reflected light it is red. After an equal volume of distilled water has been added, it becomes translucent, and viewed by reflected light it is of a dark lake colour. The difference is owing to the action of water on the red corpuscles ; the hæmoglobin has been dissolved, and,

leaving the stroma, has become diffused throughout the fluid ; the colourless stromata permit the transmission of light through the fluid, and less light is reflected from it. Hence it is translucent when viewed by transmitted light, dark lake-coloured when viewed by reflected light. Blood may be 'laked' by other means than water—*e.g.* by alternate freezing and thawing, or by the addition of chloroform.

**The composition of plasma, lymph, chyle, and serous fluids.**—The proteids of blood are for the most part contained in the plasma, the corpuscles containing a comparatively small quantity of proteid, though their chief constituent, hæmoglobin, has an elementary composition closely resembling that of proteids, and yields proteid (globin) when it is decomposed by heat or by acids. The proteids of the *plasma* are fibrinogen, paraglobulin, and serum-albumin ; the last two are in about equal, but somewhat variable proportion, and amount to between 7 and 8 per cent. of the liquid ; fibrinogen is present in very small amount, less than 1 per cent. *Serum* differs from plasma in that it contains no fibrinogen ; it contains paraglobulin and serum-albumin in the same proportion as does the plasma. Thus it appears that these fluids are, above all, highly albuminous solutions, containing nearly as much proteid as is contained in ordinary white of egg. Normal *lymph* is essentially a dilute plasma ; it is, like it, a coagulable albuminous solution with roughly 5 per cent. of proteids—namely, fibrinogen, paraglobulin, and serum-albumin. *Chyle* is lymph plus fat, and its composition differs with the state of digestion ; chyle taken from an animal in full digestion of fat has been found to contain as much as 15 per cent. of fat, *i.e.* three times as much as is contained in ordinary milk. All the above-named fluids, with the exception of serum, are spontaneously coagulable ; plasma yields the firmest and most abundant clot, which is, in fact, a blood-clot minus the red corpuscles, and the fluid in which the clot floats is serum ; lymph, being a more dilute fluid, yields a less considerable clot, and chyle yields a soft white clot like a loose curd.

Abnormal lymph, such as gathers in serous sacs, constituting *serous effusions* (pericardial, pleural, peritoneal and hydrocele fluids), varies as regards its proteids and as regards its coagulability. All serous effusions contain proteids (serum-albumin and serum-globulin) in proportions varying between .5 and 5 per cent. Some serous effusions coagulate spontaneously ; others only after



addition of serum or of fibrin-ferment; others, again, are quite uncoagulable. An effusion of the first kind amounts practically to a dilute lymph, and contains, in addition to the serum-proteids, fibrinogen, fibrin-ferment, and a considerable number of leucocytes: an effusion of this kind has usually formed rapidly in consequence of an acute inflammation of a serous membrane. Effusions of the third kind amount practically to a dilute serum; they are usually chronic, and contain only a small proportion of proteids with little or no fibrinogen and no ferment. Effusions of the second class are intermediate between the two extremes; they contain fibrinogen as well as the two serum-proteids, but no fibrin-ferment; they do not coagulate spontaneously, but only after the addition of blood-serum or shreds of fibrin—in short, of any substance containing fibrin-ferment. The relations that these effusions bear to plasma and lymph on the one hand, and to serum on the other, are exhibited in the following table:—

Plasma and Lymph	Serous effusions			Serum
Coagulable	I. Coagulable	II. Coagulable by ferment	III. Uncoagulable	Uncoagulable
Serum-albumin Serum-globulin Fibrinogen Fibrin-ferment	Serum-albumin Serum-globulin Fibrinogen Fibrin-ferment	Serum-albumin Serum-globulin Fibrinogen —	Serum-albumin Serum-globulin — —	Serum-albumin Serum-globulin — Fibrin-ferment

As regards differences of composition in the lymph coming from various different sources there is little definite to be said. The sub-arachnoid or cerebro-spinal fluid (p. 522), which is in fact the lymph of the central nervous system, is peculiar in that it contains a reducing body—formerly assumed to be sugar, but now recognised to be an aromatic compound—pyrocatechin.

**Facts and theories relating to coagulation.**—It is with the second variety of serous effusion—usually pericardial fluid or the fluid of hydrocele—that certain classical experiments have been made bearing upon the theory of coagulation. Buchanan, in 1831, and Alexander Schmidt, in 1861, observed that such a serous fluid—*e.g.* hydrocele fluid—uncoagulable by itself, yielded a clot if mixed with serum and left to stand.

It was supposed by Buchanan that a substance dissolved in the serum effected the transformation into fibrin of a substance dissolved in the hydrocele or other fluid. He found that the



same transformation could be effected by washed blood-clot, by portions of buffy coat, &c., and that the efficiency and rapidity of clotting were proportionate to the quantity of white corpuscles present. From which he concluded that the transforming agent was derived from the white corpuscles, and he compared its action to that of rennet upon milk. Translated into modern language, Buchanan's explanation of the experiment would run thus : a *ferment* dissolved in serum and derived from the leucocytes effects the transformation into fibrin of *fibrinogen* dissolved in the hydrocele or other fluid. In the experiments above described, the two fibrin factors are separately present each in one of two liquids which give a clot when they are mixed ; in the case of the blood or blood-plasma, both factors are present in the plasma, and the blood or blood-plasma is of itself coagulable.

A. Schmidt separated and named the proteids of these various fluids ; from hydrocele fluid he separated the fibrin antecedent, and gave to it its name, fibrinogen ; he found that the fluid after separation of fibrinogen was no longer coagulable with serum, while the separated fibrinogen dissolved in saline solution, and, mixed with serum, might coagulate ; from serum he separated a body which—on account of its presumed fibrin-forming effect—he named *fibrinoplastin* ;<sup>1</sup> he regarded this fibrinoplastic body as a product of disintegrated leucocytes, and he showed that a solution of this body acts upon hydrocele fluid or fibrinogen solution in the same manner as ordinary serum. But he found that it was possible for these two bodies to be present together in the same liquid, *e.g.* hydrocele fluid, which does not coagulate until serum is added. He concluded that serum contains a ferment, also derived from the leucocytes, and that the union of the two globulins was brought about by the presence of this ferment. Schmidt's theory assigns the fibrin-formation in coagulation to three fibrin-factors : (a) fibrinogen, (b) fibrinoplastin, (c) fibrin-ferment.

The researches of Hammarsten have gone to show that paraglobulin is not an indispensable factor in coagulation, and bring us back to the view that coagulation depends upon the conversion into fibrin of a single body (fibrinogen) under the influence of a ferment (fibrin-ferment) which is derived from the leucocytes. Paraglobulin plays only an accessory part in the process ; its

<sup>1</sup> 'Paraglobulin' and 'serum-globulin' are synonymous terms, denoting the same body ; Schmidt's 'fibrinoplastin' forms part of, and is indistinguishable from, the serum-globulin.

presence leads to a more decided and abundant formation of fibrin, an influence which is possessed by many other bodies—by casein, by calcium salts, and by nitrogenous derivatives of the proteids, such as lecithin, glycin, leucin, tyrosin, &c.

Among the bodies found to be adjuvant of coagulation, the calcium salts are the most remarkable; in the absence of calcium no coagulum is formed, either in blood by 'fibrin-ferment,' or in milk by 'rennet-ferment,' or in muscle by 'myosin-ferment,' and in each case the body when forming a coagulum appears to include calcium in its molecule. Oxalates abolish the coagulability of blood (or of milk or of muscle-plasma) by combining with its calcium salts; a mixture of 9 parts of fresh blood with 1 part of a 1 per cent. solution of potassium oxalate is thus 'decalcified,' and remains fluid until it is recalcified (Arthus and Pagès).

The tendency of the present day is to assign the essential fibrinoplastic action to nucleo-albumin, and a lively controversy has centred itself round the leucocyte. Is it identical with a lymph-cell? Does it disintegrate in shed blood? Is it a source of fibrin-ferment, &c., &c.? Wooldridge pointed out the fibrino-plastic action, first of lecithin, then of tissue-fibrinogens, and, finally, gave fibrin to be the resultant of two plasma-globulins, termed by him 'A fibrinogen' and 'B fibrinogen.' Halliburton regards fibrin-ferment as a globulin, refers to tissue-fibrinogen as cell-globulin, and, admitting the association of nucleo-albumin with cell-globulin, considers nucleo-albumin to be the essential fibrinoplast. He finds that the fibrinoplastic action of serum goes with the precipitate (globulin and nucleo-albumin) produced by saturation with magnesium sulphate. Schmidt holds to the leucocyte, and asserts that it disintegrates in shed blood; he admits that an anti-fibrinoplastic, as well as a fibrinoplastic body, are thus produced; he finds that an alcoholic extract of lymph-gland is fibrinoplastic, and that an aqueous extract of the residue is anti-fibrinoplastic. Lilienfeld goes further in the same direction; according to him the two bodies occur, combined in normal leucocytes, as a nucleo-albumin, which, under the influence of calcium salts, divides into nuclein, a promoter of coagulation, and an albumose body ('histone'), a restrainer of coagulation.

The chief points of unanimity in the experimental debate are to the following effect: The substances governing fibrin-formation in the living body are derived from tissue-cells (*e.g.* of lymph-glands); these cell-derivatives may be favourable or unfavourable to coagulation—fibrinoplastic or anti-fibrinoplastic; the former contain much phosphorus and are allied to nuclein, the latter are allied to the albumose group. The action of an indefinite fibrin-ferment is now assigned to the presence of the somewhat less indefinite body 'nuclein,' of which we know that it contains a high percentage of phosphorus.



**Methods of separating the proteids of plasma and of serum.**—*Fibrinogen* may be prepared from ‘salted plasma,’ or from pericardial or hydrocele fluids, and the best method is that known as the sodium chloride method (Hammarsten), the main purpose of which is the separation of fibrinogen from paraglobulin by virtue of the differing solubilities of these bodies in sodium chloride solutions. Fibrinogen is insoluble in a semi-saturated solution, and is therefore precipitated by adding to the plasma an equal volume of a saturated solution; whereas paraglobulin is not thus precipitated, being soluble in a semi-saturated solution. The precipitated fibrinogen is redissolved in a quarter-saturated solution of sodium chloride, and re-precipitated by a saturated solution; this process is repeated three times, in order to purify the final product, and the residual salt is got rid of by dialysis. ‘Salted plasma’ is obtained by centrifuging a mixture of three volumes of blood drawn off from an artery into one volume of a saturated solution of magnesium sulphate; it is the best fluid to use for the present purpose, as the fibrinogen obtained from serous fluids is apt to include lecithin, however much it be purified.

*Paraglobulin* or *serum-globulin* is best prepared from serum. Saturation with sodium chloride (Hammarsten), or prolonged dialysis (A. Schmidt), or careful neutralisation with dilute acetic acid (Panum), or the prolonged passage of  $\text{CO}_2$  through dilute serum (A. Schmidt), or semi-saturation with ammonium sulphate, all cause a more or less complete precipitation of paraglobulin from serum. Saturation with magnesium sulphate is, however, the most approved method, and gives a complete precipitation of paraglobulin (Hammarsten). To estimate the amount of globulin present, the fluid must be completely saturated with  $\text{MgSO}_4$ , and to this end it is necessary to subject it to prolonged agitation with an excess of the salt. The precipitate is collected on a filter, washed with saturated  $\text{MgSO}_4$  solution dried at  $100^\circ$ , washed with boiling water, and finally dried and weighed.

To prepare *fibrin-ferment*, serum is mixed with twenty times its bulk of alcohol, and allowed to stand for several weeks; the proteids are thereby coagulated and form a deposit at the bottom of the flask. The alcohol is poured off and the residue extracted with water and filtered. The filtrate contains very little proteid, but possesses the same property of promoting the coagulation of, *e.g.* hydrocele fluid, which is possessed by ‘washed clot’ or

fibrin or serum, *i.e.* it contains fibrin-ferment in solution. This ferment has not, however, been isolated as a solid substance.

Blood received into alcohol directly from an artery, and the residue subsequently extracted with water, yields a solution possessing the properties of fibrin-ferment, though in much less degree than a solution prepared from serum or from blood which has been shed some time. This, according to Schmidt, may be accepted as evidence that the ferment is normally present in small proportion in circulating blood.

*Serum-albumin* is obtained as follows: paraglobulin is precipitated by the complete saturation of serum with  $\text{MgSO}_4$ ; the filtrate, which now contains only serum-albumin, is dialysed for twenty-four hours or more to remove the magnesium sulphate; the solution is evaporated *in vacuo* over sulphuric acid, or in a shallow vessel at a temperature not exceeding  $40^\circ$ . Serum-albumin is thus obtained as a yellowish, brittle solid; its solutions coagulate between  $70^\circ$  and  $85^\circ$ , according to their acidity and concentration.

The points by which solutions of fibrinogen and of paraglobulin are distinguished are (1) the influence of fibrin-ferment, (2) the effect of semi-saturation by  $\text{NaCl}$ , and (3) their temperature of coagulation. Fibrinogen comes down at  $56^\circ \text{C.}$ , paraglobulin at  $70^\circ$  to  $75^\circ$ .

**Influence of the vessels and of the tissues upon coagulability.**—Normally the blood remains fluid while it is within the vessels, and coagulates shortly after it is shed. Abnormally it may coagulate in the vessels, or it may fail to coagulate after removal from them.

1. *Normally the blood remains fluid while in the vessels, and coagulates after removal from them.*—The production of fibrin-ferment from leucocytes probably takes place continually though slowly, in circulating as well as in shed blood, and it is moreover probable that this fibrin-ferment is somehow disposed of and removed by the living vessels as fast as it is formed. This may be regarded as an explanation of the non-coagulation of blood under normal circumstances, and certain facts may be adduced to illustrate this presumed action of living vessels.

A jugular vein full of blood tied in two places and cut out of the body keeps the blood fluid for hours; the same thing may be done with a turtle's or frog's heart. If the blood be allowed to escape an hour or two after the vessel has been excised, it forthwith coagulates, but if it be left in the vessel it may remain



fluid indefinitely and begin to decompose. That this restraining action of the vessels is peculiar to intact tissue is indicated by the fact that in the body blood that escapes from a ruptured vessel forms clots, although it is in contact with living tissues. Serous effusions (of the first class alluded to above) are likewise kept fluid by the influence of the living membrane enclosing them; the pericardial fluid of the horse, for instance, coagulates spontaneously if it is removed immediately after death, but does not do so if it is left for a few hours undisturbed in the pericardium, *i.e.* under the influence of a living membrane.

2. *Abnormally the blood may coagulate in the vessels, or may fail to coagulate after removal from them.*—The blood is liable to coagulate in an injured or diseased vessel, and it sometimes happens that the blood coagulates *en masse* in a large area of uninjured vessels—in the pulmonary or in the portal vessels, for instance—causing sudden death by complete arrest of the circulation. The local coagulation is brought about by the gradual deposition of fibrin upon the roughened or irregular internal surface of a vessel, preceded and, in all probability, caused by an accumulation of leucocytes and of blood-platelets, which adhere to the rough surface and disintegrate; the resulting clot is called a ‘thrombus,’ and the process of coagulation under these circumstances is termed ‘thrombosis.’ General coagulation may be experimentally effected by the injection of putrid matter, or of hæmoglobin, or of lymph, or of extracts of the thymus gland and of the testis. According to Schmidt, it is due to an excessive production of fibrin-ferment, but to this view it may be objected that large injections of the latter fail to produce coagulation; according to Wooldridge, it is due to the tissue-fibrinogens alluded to above. It is one of the dangers attendant upon transfusion.

Peptones—or, more accurately speaking, albumoses—injected in suitable quantity into the blood-vessels of a dog ( $\cdot 3$  gramme per kilogramme body-weight) render the blood temporarily uncoagulable; blood taken from the animal directly or within an hour after injection fails to coagulate, and its leucocytes are greatly reduced in number (Wright). Little by little coagulability is restored: blood drawn a few hours later, or on the following day, coagulates normally; the ‘peptone’ has been eliminated through the kidneys. It is to be observed that the effects entirely fail on the rabbit, and are not so well marked on cats as they are on dogs. The abolition of coagulability is associated with a con-



siderable fall of blood-pressure due to dilatation of the splanchnic vessels, and by a great diminution of  $\text{CO}_2$  in the blood. The addition of decomposing albumen, and the passage of carbon dioxide cause peptone-blood to coagulate; it may also coagulate when decomposition sets in. Similar effects on circulating blood have been produced by the injection of leech-extract, and the coagulation of freshly drawn blood can be arrested by leech-extract (Haycraft), which owes this property to the presence of an albumose (Dickinson).

Blood made to pass repeatedly through the heart and lungs, and kept away from the systemic circuit, gradually loses its coagulability (Ludwig and Pawlow); and according to Bohr, it is sufficient to prevent blood from passing through the portal system to cause coagulability to be lost.

**The life-history of the blood-corpuscles.**—The *red corpuscles* make their first appearance in the embryo as the daughter cells of branched connective-tissue cells which become modified into vessels. These *embryonic corpuscles* are nucleated, they multiply by cell-division, and give rise to non-nucleated corpuscles of the adult form. In the human embryo, up to the end of the first month, all the red corpuscles are nucleated; by the end of the third month, only 10 to 20 per cent. of their number are nucleated; at birth all are non-nucleated, as in the adult. The supply of red corpuscles is kept up in the adult (1) from the *red marrow of bone*, from which arise nucleated red corpuscles (Neumann's cells) resembling those of the embryo; (2) from the leucocytes or lymph-corpuscles, which are themselves derived from lymphatic glands and tissue, and from the spleen; (3) from the so-called blood-platelets (Bizzozero), or hæmatoblasts (Hayem), and from elementary particles (Zimmermann), the derivation of all of which is obscure. Of these various sources that first named, viz. the marrow of bone, is the most important and best attested; the derivation of red corpuscles from leucocytes, &c., is a supposition rather than an observed fact. The chief known or surmised seats of the destruction of red corpuscles in the adult are the spleen and liver; their length of life has been estimated to be from two to three weeks, the estimate, which is necessarily very rough, being based upon a comparison of the amount of pigment per diem in bile, urine, and fæces, with the total amount of hæmoglobin from which these pigments are derived.

The *white corpuscles* arise from lymphoid tissues generally

(lymphatic glands, adenoid tissue, spleen), where they multiply by cell-division. They exhibit several fundamental life-characters, and several definite observations have been made which show that they play an important part in growth, in absorption, in disintegration, and in repair. They are excitable and contractile, they can move spontaneously by slow protrusions and retractions, in a manner quite similar to that in which amoebæ are observed to alter shape and change position, hence their movements are termed *amœboid*. They can surround particles of matter with which they come in contact, and carry them as they move from place to place. They may attach themselves to the walls of the minute veins, penetrating them, and escaping into the surrounding tissues (*emigration* or *diapedesis*), where they may accumulate and form an abscess, or become organised

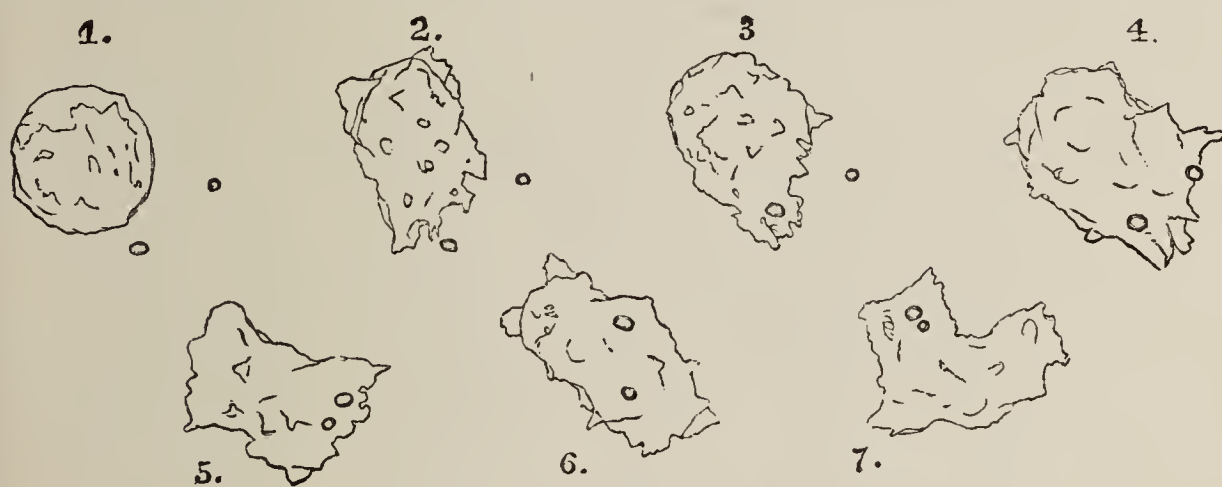


FIG. 6.—AMŒBOID MOVEMENTS.

Changes of form of a white corpuscle of newt's blood. Ingestion of two small starch granules, and their changes of position within the corpuscle.—Quain's 'Anatomy.'

and form new tissue. They are possibly agents in the absorption of fat, acting as carriers from the epithelial surface of the intestine to the lacteals of the villi; in the lungs they are certainly carriers of foreign particles that may have been inhaled; it is by their agency that particles of inhaled coal-dust may be conveyed to, and deposited in, the pulmonary lymphatic glands; in the intestines, and elsewhere, they 'englobe' and destroy deleterious microbes; it is in view of this that such leucocytes have received the name of 'phagocytes.'<sup>1</sup>

Of the various facts above enumerated, that which is of widest significance is the *emigration of leucocytes*. When there is irritation of a part there is a determination of blood to that part ('ubi stimulus ibi affluxus'), the vessels dilate and become congested, and the leucocytes in particular adhere to the vessel-walls,

<sup>1</sup> φάγειν, to eat; λευκός, white; κύτος, a cell.



accumulate there, and begin to emigrate. With excessive irritation or peculiar debility of the part, the change exceeds congestion, and is termed inflammation; with inflammation of sufficient severity, or with tissues of unusual debility, the emigrant cells may overmaster and destroy tissue, gathering into large accumulations of cells that are now called pus-cells, and forming abscesses, the contents of which must be evacuated. A pimple that comes to a head illustrates this process on a small scale; the 'matter' that escapes when the pimple bursts or is pricked, is composed of pus-cells which are emigrated leucocytes, indistinguishable from them under the microscope, and exhibiting, like them, amœboid movements.

Emigrated leucocytes may, however, play another part and effect repair of tissues: a clean, healthy wound becomes in a few



FIG. 7.—EMIGRATION OF LEUCOCYTES.

Vessels of the inferior surface of the frog's tongue as they appear after the escape of the corpuscles, filled with stationary blood, deformed and indented at the points of escape, near which the corpuscles are generally found. A portion of a vessel with an internal current is likewise seen with discs and internal and external corpuscles.—After Waller. *Phil. Mag.* 1846. 'Microscopic Observations on the perforation of the Capillaries by the Corpuscles of the Blood.'

hours covered with a layer of lymph, which is mainly composed of emigrated leucocytes. This exudation of lymph is the first step towards healing; if repair progresses in a healthy manner, the leucocytes may become organised and converted into connective tissue, forming the cicatrix by which the wound is closed.

Finally, we recognise as possible and probable (though the

fact is not proved by direct observation) that emigrant leucocytes play a part abnormally in the production of many tumours, as well as normally in the natural growth of tissue.

To sum up—let us imagine that several leucocytes start in life together from lymphoid tissue and get into the blood. One leucocyte may get attached to a vessel of an inflamed part, may pass through it and go to the bad as a pus-cell, the constituent of an abscess: another may escape at the surface of a wound, and settle down as a connective-tissue cell, the constituent of a cicatrix: a third may play the part of scavenger, and remove a foreign particle or destroy a microbe.

These are suppositions, but suppositions that have reasonable foundation in observed facts, of which amoeboid movement and emigration are the parent facts: leucocytes have been made to take up coloured granules, and by this means subsequently identified as pus-cells in an inflamed cornea; their behaviour towards microbes has been watched under the microscope; they have been found crowding round foreign bodies that are becoming absorbed or surrounded by a false membrane. But the actual transformation of leucocytes into connective tissue is not an observed fact—the origin of a tumour-cell, or of a normal tissue-cell from emigrated leucocytes has been imagined but not seen, and although the idea may be true, these facts have not yet been verified.

*Quinine* exercises a remarkable influence upon the movements of leucocytes; applied in dilute solution ( $\frac{1}{1500}$ ) to a slide of newt's blood, the amoeboid movements of leucocytes are arrested; applied locally to a frog's mesentery, or injected into the blood-vessels, it arrests emigration of leucocytes (Binz).

**The blood-gases.**—100 c.c. of blood contain between 50 and 60 c.c. of the gases *oxygen* and *carbon dioxide*. These gases are present in small proportion simply dissolved, but for the most part in a state of loose chemical combination, from which they can be separated by subjecting the blood to a vacuum. Oxygen is held by the red corpuscles in combination with hæmoglobin, as oxyhæmoglobin; carbon dioxide is held by the plasma in combination with soda, as sodium carbonate ( $\text{Na}_2\text{CO}_3$ ) and sodium bicarbonate ( $\text{NaHCO}_3$ ). In both cases the combination is very feeble; it is easily unmade, and as easily re-made. By simply subjecting blood to a vacuum, its gases, relieved from atmospheric pressure, are disengaged and may be collected, and its hæmoglobin becomes 'reduced.' Or by adding a 'reducing



agent'<sup>1</sup> to the blood, oxygen is withdrawn from combination with hæmoglobin, which remains therefore as reduced hæmoglobin.

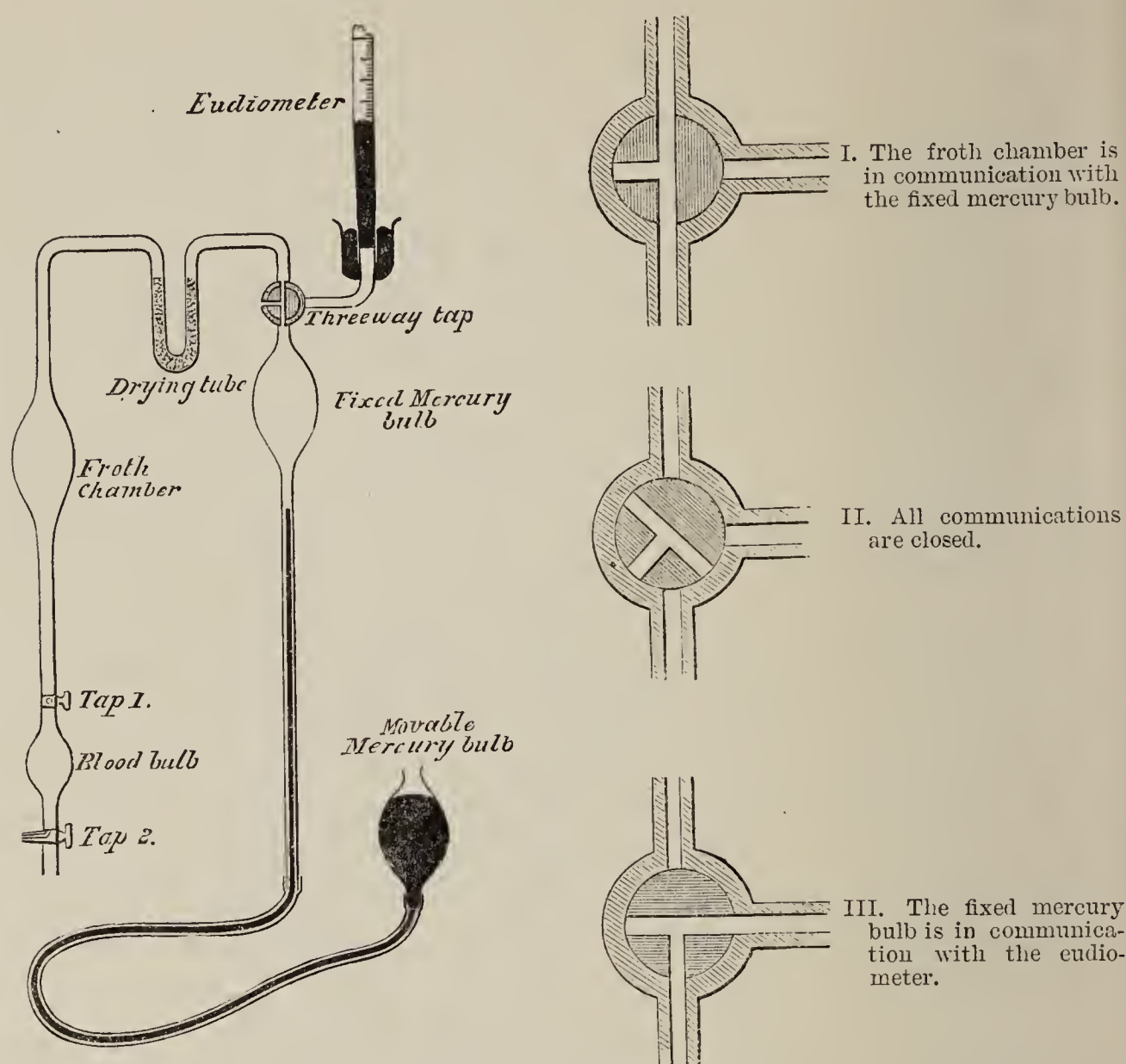


FIG. 8.—PLAN OF MERCURY PUMP FOR THE EXTRACTION OF THE GASES OF THE BLOOD.

Two glass bulbs containing mercury; one is fixed, the other can be raised and lowered. A three-way tap, which can be turned into three positions, I. II. III. A 'blood bulb,' which is filled from an artery or vein, through a tube slipped over the end of tap 2, and which can be connected with, or disconnected from, the froth chamber by opening or shutting tap 1. 'Froth chamber' serving to intercept spurts of blood when the gases are bubbling out. Vacuum is established by repeatedly lowering the movable mercury bulb while the three-way tap is in position I. and raising it with the tap in position III. The first step draws air from the apparatus into the fixed mercury bulb, the second step drives out this air by the delivery tube.

Vacuum having been established, the eudiometer filled with mercury is adjusted over the delivery tube, and the tap between froth chamber and blood bulb is opened. The blood begins to bubble and its gases are given off into the vacuum; by raising the mercury bulb (with tap as in III.) the blood-gas is discharged through the delivery tube into the eudiometer, by lowering the bulb (with tap as in I.) the extraction is repeated; by further repetition of the up and down manœuvre, and by immersing the blood bulb in water warmed to 40°, all the gas is extracted and collected in the eudiometer for further examination.

<sup>1</sup> A reducing agent in relation to blood is any compound or element having an affinity for oxygen greater than that of hæmoglobin: ammonium sulphide and Stokes's fluid are the reducing agents commonly employed. Stokes's fluid is



Germinating yeast, putrefactive bacteria, and living tissues are also reducing agents, inasmuch as they require oxygen, and will withdraw it from oxyhæmoglobin. A little yeast mixed with arterial blood and kept warm in a stoppered bottle will quickly effect the reduction ; decomposing blood—*i.e.* blood in which putrefactive bacteria are living and multiplying—blackens and becomes reduced ; and the living tissues change the blood which passes through them from the arterial or oxygenated, to the venous or deoxygenated state.

If arterial blood be left to itself in a well-stoppered bottle it will darken in colour, and become reduced before the occurrence

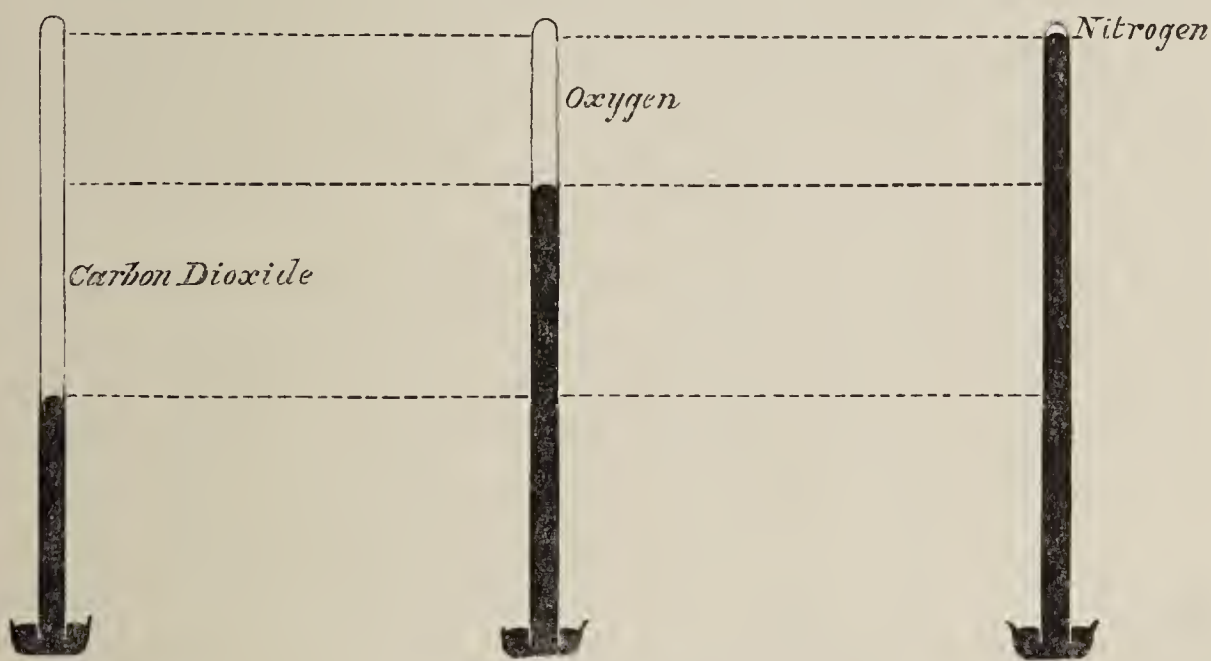


FIG. 9.—EUDIOMETER TUBES.

The gases thus obtained consist of carbon dioxide, oxygen, and nitrogen. The amount of carbon dioxide is ascertained by introducing some caustic potash into the eudiometer ; the difference in the volumes of gas in the tube before and after the absorption of carbon dioxide makes known the amount of this gas. The amount of oxygen is similarly ascertained by observing the diminution of volume consequent upon the introduction of phosphorus or of pyrogallie acid. The residual gas in the tube is nitrogen. The readings must be corrected for temperature and pressure.

of decomposition. Blood removed from the body does not die at once, nor is it wholly dead even after coagulation has taken place. As it continues to live, so it continues to consume oxygen, and, preserved from contact with the air, it consumes its own oxygen. One kilo of fresh blood will consume about 30 c.c. of oxygen per hour, as compared with one kilo of freshly excised muscle, which consumes about 300 c.c. in the same space of time.

**Hæmoglobin—its compounds and derivatives.**—Hæmoglobin—the most important constituent of red blood-corpuscles—is a prepared as follows:—To a fresh solution of ferrous sulphate, 2 grammes per 100 c.c. add 3 grammes tartaric acid ; to this solution add ammonia until it is faintly alkaline.

crystalline body, but although crystalline it is indiffusible; it closely resembles a proteid in chemical constitution, but is peculiar in that its molecule contains iron (about  $\cdot 5$  per cent.), in addition to the ordinary proteid constituents, C.O.H.N.

To obtain crystals of hæmoglobin, blood is 'laked,' *i.e.* its hæmoglobin is made to pass into solution, either by treatment with water, or with chloroform, or by alternate freezing and thawing; crystals are formed as this solution evaporates. But these crystals are not obtained with the same facility from all animals, nor of the same shape; they are easily obtained from the rat, mouse, guinea-pig and squirrel, less easily from dog and man, horse and sheep, less easily still from frog's blood.

Crystallised hæmoglobin readily absorbs and combines with oxygen, and will give off oxygen when exposed to a vacuum; at normal atmospheric pressure 1 gramme hæmoglobin absorbs 1.7 c.c. oxygen.

The two states of hæmoglobin in the blood, as oxyhæmoglobin and reduced hæmoglobin, have been already referred to. They are its normal physiological states in the red corpuscles, which are its vehicle. Hæmoglobin may, by accident or by experiment, be made to enter into two other combinations—with carbon

monoxide, forming carboxyhæmoglobin, and with nitric oxide, forming nitroxyhæmoglobin. Both these combinations are more stable than oxyhæmoglobin; oxygen is easily separated from hæmoglobin, carbon monoxide is separated with greater difficulty, nitric oxide with still greater difficulty. By passing a current of carbon monoxide through a solution of oxyhæmoglobin, oxygen is quickly and easily displaced, and carbon monoxide takes its place volume for volume in combination with hæmoglobin.

A stream of CO, or of coal-gas passed through blood, or through a solution of hæmoglobin, gives a bright red colour, which is not altered by reducing agents, while the colour of oxygenated blood is markedly altered by such

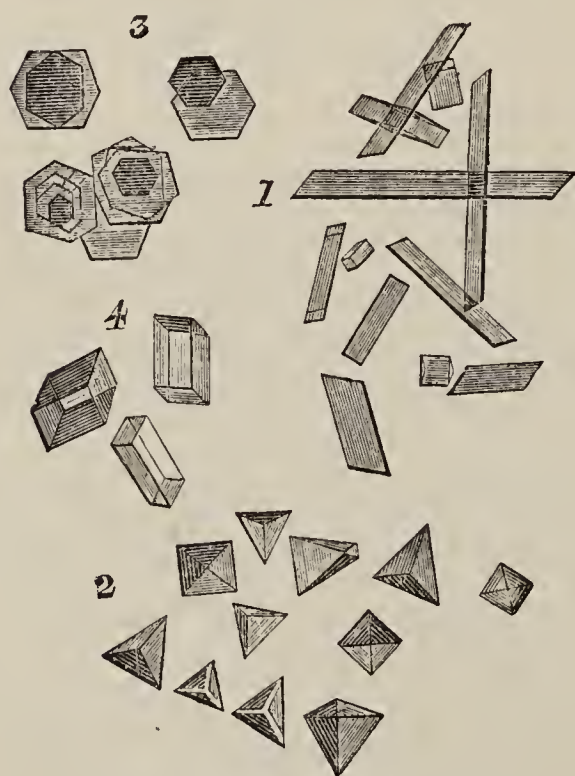


FIG. 10.—HÆMOGLOBIN CRYSTALS.

1. Man or rat. 2. Guinea-pig.  
3. Squirrel. 4. Hamster.—(Quain's  
'Anatomy.')



agents. By passing a current of nitric oxide through a solution of carboxyhæmoglobin (in the absence of oxygen), carbonic oxide is expelled, and the combination nitroxyhæmoglobin formed.

Carboxyhæmoglobin may be formed accidentally as well as experimentally. Common coal-gas contains nearly 10 per cent. of carbon monoxide; the imperfect combustion of carbon—as in charcoal stoves, or as sometimes happens in coal-mine explosions (Haldane)—entails a production of carbon monoxide, and the respiration of an atmosphere charged with carbon monoxide in any of these ways may be fatal to life, carbon monoxide taking the place of oxygen in combination with hæmoglobin, and annulling its respiratory function. The comparative stability of the compound renders this mode of poisoning all the more dangerous, and decreases the chance of recovery; still, it is important to remember that by the prolonged passage of oxygen through a solution of carboxyhæmoglobin the oxygenated state can be recovered, so that in the case of a person whose blood has been more or less profoundly poisoned, artificial respiration should be vigorously persevered in. On the other hand, the stability of the compound facilitates the recognition, in doubtful cases, of the cause of death.

The spectrum of carboxyhæmoglobin, or of the blood of a person poisoned by inhalation of carbonic oxide, is hardly distinguishable from that of oxyhæmoglobin, even when both spectra are simultaneously observed close above each other. But the addition of a reducing agent, such as ammonium sulphide, at once establishes the difference. Oxyhæmoglobin is reduced, and its two-banded spectrum disappears, being replaced by the single-banded spectrum of reduced hæmoglobin. On shaking the solution with air, re-oxygenation takes place, and the two-banded spectrum reappears; the latter soon disappears again if the solution is left standing, and if the reducing agent has been added in excess. Carboxyhæmoglobin suffers no such change on the addition of reducing agents; its two-banded spectrum remains unaltered.

The *derivatives* of hæmoglobin are bodies which are produced by its decomposition; some are normally formed in the human body, others occur abnormally in the body, and can be artificially produced out of the body.

The normal derivatives are the *bile and urine pigments*. The abnormal derivatives are *hæmatoidin*, *methæmoglobin*, *hæmatin*, and *hæmatoporphyrin*. The artificial derivatives are *hæmochro-*

*mogen*, *hæmatoporphyrin*, and *hæmin* : methæmoglobin and hæmatin can also be artificially produced from hæmoglobin.

With regard to the natural derivatives, little is known beyond the fact that the pigments of the bile and of urine (bilirubin, urobilin) are derived from hæmoglobin. *Hæmatoidin* is a crystalline substance found in old blood-clots in the brain and other organs ; it is evidently derived from hæmoglobin, but it does not contain iron, and it gives the same reactions as *hydro-bilirubin*,

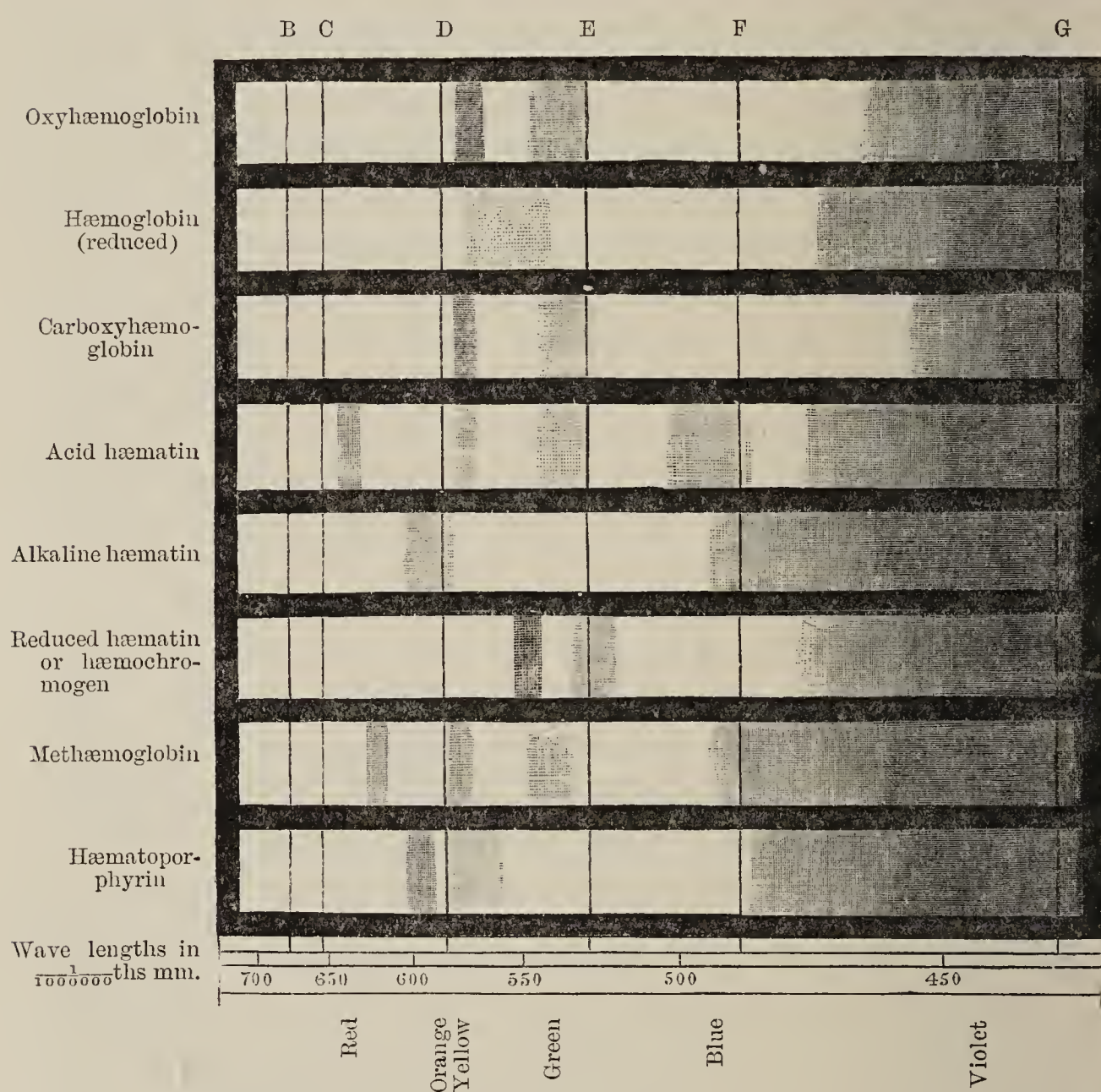


FIG. 11.—BLOOD SPECTRA.

with which it is therefore regarded as identical. Hæmatoidin thus furnishes accidental evidence of the derivation of bilirubin from hæmoglobin. *Methæmoglobin* and *hæmatin* are met with in pathological products—in ‘smoky’ urine, *i.e.* urine containing blood which has transuded from the kidney—and in ‘coffee-ground’ vomit, *i.e.* more or less digested blood from bleeding vessels in the stomach. Methæmoglobin is simply a modified kind of oxyhæmoglobin ; hæmatin is one of the products of the



decomposition of hæmoglobin. The two bodies give very similar spectra, but may, however, be easily distinguished by the effects of a reducing agent, such as ammonium sulphide : the spectrum of methæmoglobin becomes altered and gives place to that of reduced hæmoglobin, which, when the fluid is aërated by shaking, is replaced by the spectrum of oxyhæmoglobin ; the spectrum of hæmatin is also altered by the reducing agent, and gives place to that of reduced hæmatin, on re-oxygenation the spectrum of hæmatin reappears. Thus the chief distinction between the spectra of the two bodies is that from methæmoglobin the spectrum of oxyhæmoglobin can be recovered, while from hæmatin it cannot. The blood of animals that have been poisoned by inhalation of nitrite of amyl has a peculiar chocolate-brown colour, and gives the spectrum of methæmoglobin. Both these bodies may also be classed as artificial derivatives, although, perhaps, methæmoglobin should be regarded as modified oxyhæmoglobin rather than as a decomposition product. By adding acetic acid to diluted blood, the hæmoglobin is decomposed into hæmatin and a proteid ; by shaking this fluid with ether a coloured ethereal solution is obtained, which gives the spectrum of *hæmatin in acid solution*. On rendering this fluid alkaline by the addition of caustic potash in excess, the spectrum is altered to that of *hæmatin in alkaline solution*. On adding a reducing agent to the alkaline solution the spectrum is altered to that of *reduced hæmatin*. If the blood be decomposed by caustic potash in the absence of oxygen, a body called *hæmochromogen* is produced, which is identical with reduced hæmatin, giving the same spectrum, and, as soon as it comes in contact with air, giving the spectrum of hæmatin. Hæmatin contains all the iron of hæmoglobin ; if, however, hæmoglobin be decomposed by still more powerful means than above mentioned—by strong sulphuric acid, or by heating with fuming hydrochloric acid—a more profound decomposition is effected, and as the result a body called *hæmatoporphyrin* is obtained, the iron being separated by the acid in the form of ferrous sulphate or chloride.

*Hæmin* is a compound of hæmatin with hydrochloric acid. It is a crystalline body of great importance in medico-legal practice, as it furnishes a most reliable test of the nature of stains supposed to be of blood. By boiling on a slip of glass a little dried blood with glacial acetic acid and a minute fragment of sodium chloride, this compound is formed, and can easily be recognised

under the microscope as small brown rhombic crystals. They are commonly spoken of as Teichmann's crystals.

To sum up—we have learned that hæmoglobin can combine with  $O_2$  and with CO in or out of the body; that in the body it may decompose and give rise to hæmatin or to the iron-free body, hæmatoidin or bilirubin; that analogous changes can be brought about outside the body by the action of reagents, hæmatin being produced by comparatively weak reagents, and an iron-free body, hæmatoporphyrin, by strong acids. Methæmoglobin is so little removed from oxyhæmoglobin as hardly to deserve a place among derivatives; it is produced by the weakest reagents, and it can be easily restored to the original state as oxyhæmoglobin. It is to be remembered that in the contents of the stomach and of the urinary bladder, blood-pigment may be found, in the form of oxyhæmoglobin or of methæmoglobin or of hæmatin; in one or two cases the spectrum of hæmatoporphyrin has been identified.

*Estimation of hæmoglobin.*—If 1 c.c. of normal blood and 1 c.c. of 'poor' blood be each diluted with 100 c.c. of water, and the two solutions compared, that of normal blood will be evidently of a deeper tint than that of 'poor' blood, *i.e.* the latter evidently contains less hæmoglobin. If, instead of adding at once 100 c.c. of water to 1 c.c. of poor blood, the water is added gradually until the tint is judged to be equal to that of the diluted normal blood, then from the quantity of water added we may estimate how much hæmoglobin is present in the poor blood as compared with normal blood. For instance, equality of tint being established with 20 c.c., the hæmoglobin value is  $\frac{2.0}{10.0}$  of the normal; if with 30 c.c., it is  $\frac{3.0}{10.0}$  of the normal. In the practical application of the method, smaller quantities of blood are taken, *i.e.*  $\frac{1}{50}$  c.c. = 20 cubic mm., and this is diluted until it is equal in tint to a permanent standard colour made with glycerine and carmine, and equivalent to normal blood diluted 100 times. Thus arranged, the apparatus is termed a *hæmoglobino-meter*. If the 20 cubic millimeters of blood give equality of tint with 2 cubic centimeters, *i.e.* 2,000 cubic millimeters of water, the amount of hæmoglobin is normal; if with 1,000, the amount is half normal; if with 500, the amount is one-quarter normal.

The most convenient and accurate hæmoglobinometer is that of Oliver. Blood suitably diluted in a circular cell is compared with a series of more or less deeply coloured glasses that have been carefully adjusted so that the scale of tints corresponds



with a scale of blood percentages. Separate colour scales must be used according as observations are made by daylight or candle-light.

Individual spectra may be identified by means of a double spectro-scope as follows :—

A spectrum with *two* characteristic bands might be produced by *oxyhæmoglobin*, *carboxyhæmoglobin*, *hæmochromogen*, or *hæmatoporphyrin*.

*Oxyhæmoglobin* and *carboxyhæmoglobin* would be distinguished by adding a reducing agent. *Hæmochromogen* bands would be to the violet side, and *hæmatoporphyrin* bands to the red side of oxyhæmoglobin bands.

A spectrum with one distinct band on the red side of the D line might be due to *acid hæmatin*, *alkaline hæmatin*, *reduced hæmoglobin*, or *methæmoglobin*. A reducing agent would wipe out this band in the case of methæmoglobin or alkaline hæmatin,

giving reduced hæmoglobin or reduced hæmatin ; reduced hæmoglobin could be temporarily re-oxygenated by shaking the fluid.

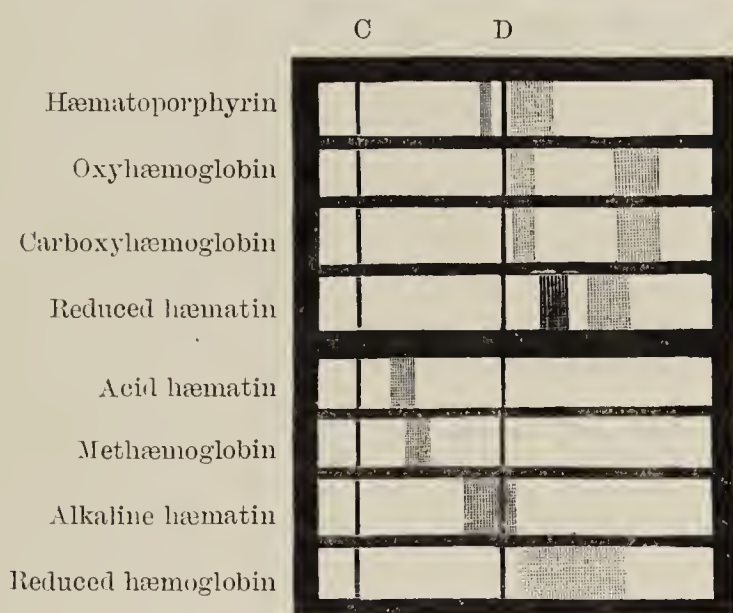


FIG. 12.

The chief difference between **arterial blood** and **venous blood** is that the former contains more oxygen and less carbonic acid than the latter ; this difference is accompanied with an evident difference of colour, arterial blood being bright red, while venous blood is almost black, and it follows that the amount of oxyhæmoglobin is greater in arterial than in venous blood, while the amount of reduced hæmoglobin is greater in venous blood. But it would be a mistake to suppose that all the hæmoglobin in arterial blood is oxyhæmoglobin, and that all the hæmoglobin in venous blood is reduced hæmoglobin. The difference is only in the relative amounts, as may be illustrated by the following example :—

100 c.c. of dog's blood from crural artery contained :

Oxyhæmoglobin	.	.	.	.	.	.	.	15 grammes
Reduced hæmoglobin	.	.	.	.	.	.	.	2 „

100 c.c. of dog's blood from crural vein contained :

Oxyhæmoglobin	.	.	.	.	.	.	.	10 grammes
Reduced hæmoglobin	.	.	.	.	.	.	.	7 „



These figures show that arterial blood is not completely saturated with oxygen, and that venous blood is not completely deoxygenated. A corresponding fact would be shown by measurements of the gases contained in arterial and in venous blood, as in the following example :—

100 c.c. arterial blood contained :

Oxygen	.	.	.	.	.	.	.	20 c.c.
Carbonic acid	.	.	.	.	.	.	.	40 „

100 c.c. venous blood contained :

Oxygen	.	.	.	.	.	.	.	10 c.c.
Carbonic acid	.	.	.	.	.	.	.	48 „

Thus arterial blood still contains carbonic acid, and venous blood still contains oxygen.

If, during life, arterial and venous blood be removed without bringing them in contact with air, and examined by the spectroscope, both will show the two-banded spectrum of oxyhæmoglobin. If, immediately after death, arterial and venous blood—*e.g.* from the aorta and from the right auricle—be similarly examined, the former will show the two-banded spectrum of oxyhæmoglobin, the latter the one-banded spectrum of reduced hæmoglobin. The tissues continue to live for some time after an animal has ceased to move and appears dead ; internal respiration by the tissues having continued after external respiration in the lungs has ceased, venous blood has become completely reduced, and is found so in the right auricle.

It is even possible to observe the reduction of oxyhæmoglobin by the living tissues of the human body. If sunlight be reflected from a finger-nail where it is pink with blood, the two-banded spectrum of oxyhæmoglobin will be visible. If a ligature be tied round the finger so as to obstruct its blood-supply, these bands will vanish, and under very favourable circumstances the dim band of reduced hæmoglobin may become visible.

Whereas arterial blood is of uniform composition, the same kind of blood being distributed to all parts of the body, venous blood, returning from different parts and organs, is not uniform, but varies in accordance with the material that may have been added to or subtracted from the blood. The blood of the *hepatic vein*, coming from the liver, where sugar is constantly produced, contains a little more sugar than other blood. The blood of the *portal vein*, coming from the digestive viscera, contains products of digestion—if digestion is taking place—an excess of sugar or

of fat or of proteid, if these substances are being absorbed from the intestinal tract. The venous blood of *secreting glands* is also apt to vary from the usual quality; from a gland that is in constant action, like the kidney, the blood returns in a less deoxygenated state than from a muscle, or from the brain. The blood of the *renal veins* is not like ordinary venous blood, but bright red, like arterial blood; and, in correspondence with the fact that the kidney separates urea from the blood, its venous blood contains less urea than other venous or than arterial blood. Other glands which are sometimes in action, sometimes at rest, show corresponding differences: the venous blood of the *sub-maxillary gland*, which is of the usual venous quality while the gland is at rest, becomes bright red, like arterial blood, while the gland is actively secreting. With *muscles* the reverse is the case: under ordinary conditions their venous blood is of the ordinary typical venous quality, but if it is analysed for gases during muscular activity, it is found to be more venous than before, and both the deficit of oxygen and the excess of carbonic acid are greater than in the blood coming from quiet muscles. From paralysed muscles the blood is, on the contrary, less venous, and both the deficit of oxygen and the excess of carbonic acid are less than before. Differences of *temperature* also obtain in different kinds of venous blood; venous blood returning in a superficial vein from an external part is cooler than the arterial blood going to the part; thus, for instance, blood in the jugular vein is of lower temperature than in the carotid artery, and the blood in the crural vein is about  $1^{\circ}$  lower than that in the crural artery.<sup>1</sup> On the other hand, deep veins leading from active organs contain blood at a higher temperature even than that of arterial blood; of this inequality the hepatic venous blood is the best known instance—its temperature is about  $1^{\circ}$  higher than that of the blood in the aorta. The blood of the *splenic vein* contains a larger proportion of leucocytes than normal blood, in consequence of the production of leucocytes taking place in the gland.

**Composition of the blood in disease.**—*Anæmia*, or poverty of the blood, is the commonest and most widespread departure from the healthy standard, and is the invariable consequence of copious bleeding or of any kind of prolonged or debilitating disease. The popular term ‘poverty of the blood’ is accurately descriptive; the blood of an anæmic person is not deficient in

<sup>1</sup> The Centigrade scale will always be referred to in the text. The relation between the Fahrenheit and Centigrade scales is given in the Appendix.



amount, but it is a weak blood, with an excess of water, and a deficiency of solids, and a low specific gravity (1030 to 1040) ; the number of red corpuscles is deficient, and the amount of hæmoglobin in these even is deficient ; contrasting the blood of a healthy person with that of a patient in extreme anæmia, there have been found, for instance :

Proportion of water . . . . .	88	instead of 80 per cent.
„ „ solids . . . . .	12	„ „ 20 „
Specific gravity . . . . .	1035	„ „ 1055 „
Number of red corpuscles per cubic mm.	1 to 2	„ „ 4 to 5 millions.
Proportion of hæmoglobin . . . . .	1 to 2	„ „ 12 per cent.

from which it is evident that the deficiency is essentially a deficiency of hæmoglobin.

*Leucocythæmia* is a state characterised by an excessive number of leucocytes in the blood, and is usually associated with excessive growth of the spleen, or of the lymphatic glands.

According to observations made in the old blood-letting days, the blood of a person suffering from severe *inflammation* (of the lung, for instance) usually yields an excessive amount of fibrin, but coagulates more slowly, with formation of the so-called buffy or inflammatory coat.

Diseases of the *kidneys* are accompanied with anæmia, and the blood contains an excess of urea. In *gout*, the function of the kidneys is generally deficient, and the blood is consequently altered as just stated, with this additional and characteristic feature, viz. excess of uric acid.

Diseases of the *liver* very commonly cause anæmia ; when the flow of bile is obstructed, bile-pigment accumulates in the blood and lymph.

The blood of patients suffering from severe *diabetes* contains an excess of sugar.

*Hæmophilia* is the name applied to a condition in which there is a peculiar tendency to bleeding, which it is difficult to check. The blood of persons subject to this affection is said to be deficient in fibrin.

**Quantity of blood in the body.**—The total amount of blood has been estimated to be in man about  $\frac{1}{13}$  of the body-weight, i.e. five to six litres (or rather more than one gallon). From this datum, and from those given on page 13, we may form a rough notion of the absolute amounts of the various proximate principles in circulation in the blood, viz. about 500 grammes hæmoglobin, 450 grammes proteids, 2·5 grammes iron, &c.



The total amount is distributed in the body approximately as follows:— $\frac{1}{4}$  in the thoracic viscera;  $\frac{1}{4}$  in the muscles;  $\frac{1}{4}$  in the liver;  $\frac{1}{4}$  in remaining parts. It is estimated by collecting all the blood that escapes when an animal is bled to death, subsequently washing out the vessels with salt solution, finally mincing and washing the organs. The amount of blood in these two washings is found by comparison of their red tint with the tint of a sample of blood of known dilution; the comparison is facilitated by the previous passage of carbonic oxide through the fluids. For example, a dog weighing ten kilogrammes is bled, and yields 400 c.c. of blood; the vessels are washed out and the various tissues are minced and washed; the united bulk of the washings is five litres, and the tint is found to be equal to that of one c.c. of blood fifty times diluted, *i.e.* the five litres contain  $\frac{1}{50}$  of blood = 100 c.c. The total amount of blood is therefore 500 c.c., that is to say,  $\frac{1}{20}$  the weight of the dog.

The estimation of the blood in particular organs is determined in a similar manner, the vessels having been previously ligatured before the cessation of circulation.

**Tests for blood.**—It is often required to determine whether or no a pathological fluid contains blood—whether or no a stain on weapons or clothes is due to blood. We have four tests by which blood may be identified. (1) By the microscope we may determine the presence or absence of blood-corpuscles in a fluid or in a fresh stain. We may be able to determine whether the blood is mammalian or that of a bird or fish, but we cannot specify what kind of mammalian animal is the source of the blood under examination. There are, it is true, differences of size between the red corpuscles of various mammalia (*vide* fig. 3), and we might thus, for instance, be able to deny that a stain caused by goat's blood was caused by human blood, but we should not be able in practice to distinguish human blood from that of animals such as the cat, dog, rabbit, ox, &c. On the other hand, we might positively distinguish the blood of fowl or fish from that of mammalia. A knife stained with blood containing oval corpuscles has, in all probability, not served to kill a man; a girl supposed to spit blood which is found to contain oval blood-corpuscles has probably obtained it from a fowl, and her statement may be disbelieved.

In many cases corpuscles may not be found and yet blood be present; it will, however, be identified by other tests, which, indeed, are in all cases to be employed as confirmatory tests. Of

these the best is (2) *the hæmin test*, applied as above stated ; the presence of hæmin crystals proves the presence of blood, but gives no information as to its kind ; any kind of blood can yield hæmin crystals, and a blood-stain, however old, will yield them.

(3) The *spectroscope* affords a convenient and expeditious means of ascertaining whether or no a coloured fluid owes its colour to blood. The fluid may contain hæmoglobin or its derivatives. An ordinary blood-stain dissolved in water gives the oxyhæmoglobin spectrum, or, if old, it may give the methæmoglobin or the hæmatin spectrum : the blood of a person poisoned by coal-gas will give the carboxyhæmoglobin spectrum : blood in the urine, or in the vomited contents of the stomach, may give either the oxyhæmoglobin, the methæmoglobin, or the hæmatin spectrum. These will be distinguished as above described, and by the alterations effected by reducing agents. It is of practical importance to remember that the spectrum of oxygenated hæmatin is much less distinct than that of reduced hæmatin ; so much so that a pathological fluid containing hæmatin may yield a blank spectrum until after the addition of ammonium sulphide. It is also to be borne in mind that the hurried examination of a pink fluid and the discovery of a two-banded spectrum are not proof positive that hæmoglobin is the colouring agent, for carmine gives a very similar spectrum. If a very small quantity of fluid is available, the spectroscope is used in combination with a microscope, being substituted for the eye-piece of the latter : an instrument of this kind is termed a *microspectroscope*.

(4) On the addition of fresh tincture of *guaiacum* and *ozonic ether* (which contains hydrogen peroxide) to a fluid which contains a trace of blood, a sapphire-blue colour is produced. The test is, however, not characteristic, for blood is not the only substance by which the colour is elicited.



## CHAPTER III

## THE CIRCULATION

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**General principles.**—The blood does not remain stagnant, nor does it ebb to and fro ; it circulates. From the heart it is driven into the arteries, from the arteries into the capillaries, from the capillaries into the veins, from the veins finally it returns into the heart. Servetus, 1553, passage of blood from right ventricle through lungs to left ventricle—Harvey, 1628, passage of blood from left ventricle through system to right ventricle—Malpighi, 1661, passage of blood from arteries through capillaries to veins. These three names and dates suffice to indicate by whom and by what stages this simple and fundamental conclusion was arrived at ; but a study of the original writings is necessary to the appreciation of the fact that Harvey discovered the ‘motion of the blood in a circle’ round lungs and system, ‘down’ arteries and ‘up’ veins, and logically demonstrated the circulation of the blood. His chief premises to the conclusion were : the disposition of valves in the veins and in the heart, the pulse in the arteries coincident with the systole of the ventricles, the great size of the minimum volume of blood that must be driven into the arteries within even a short space of time, the spurting of blood from the cardiac end of an artery, and its abundant escape from the peripheral end of a vein, and finally the effects of ‘very tight’ and of ‘moderately tight’ ligatures of the arm upon the vessels of the forearm. Harvey laid particular stress upon these last observations as demonstrating ‘that the blood enters a limb by the arteries, and returns from it by veins,’ and this portion of his argument may therefore be repeated : with a very tight ligature as employed in amputations, the arteries pulsate above the ligature, and do not pulsate beyond it ; with a moderately tight ligature, as employed in venæsection, the veins beyond the ligature are swollen and knotted, the blood in a vein can be stroked upwards, not downwards, and the vein itself will be found to refill from below ; the tight ligature stopped all circulation, the less tight ligature permitted the arterial pulse to pass, but stopped the return of venous blood, the manipulation of the swollen veins showed the direction of the blood within them, and the action of their valves. Harvey’s proof of the circulation was thus logically complete, the unseen link in his circle was seen by Malpighi with the aid of a magnifying-glass.

The heart of mammalia is a double organ, and each part of this double organ is composed of two chambers—one auricle and one ventricle ; thus the entire organ comprises four chambers, viz. the left auricle, the right auricle, the left ventricle, the right

ventricle. The arteries lead from the ventricles, the aorta or systemic artery from the left, and the pulmonary artery from the right. The veins lead to the auricles—the venæ cavæ, or systemic veins to the right, and the pulmonary veins to the left. Outside the heart the blood flows from ventricle to auricle, through arteries, capillaries, and veins; inside the heart the blood flows from auricle to ventricle. A drop of blood will complete its circulation, passing through the several chambers of the heart and through the vessels in the following order:—left

auricle, left ventricle, systemic arteries, systemic capillaries, systemic veins, right auricle, right ventricle, pulmonary arteries, pulmonary capillaries, pulmonary veins, left auricle; thus a drop of blood to complete its circulation makes two journeys from and to the heart—one through the systemic arteries, capillaries, and veins, from the left ventricle to the right au-

ricule; the other through the pulmonary arteries, capillaries, and veins, from the right ventricle to the left auricle. The first circuit is called the *systemic* or *greater circuit*; the second is called the *pulmonary* or *lesser circuit*. The first circuit conveys the blood through the whole system to all the tissues of the body; the second circuit conveys the blood through the lungs.

The force by which the blood is made to circulate is supplied by the heart muscle, which by its forcible contraction expels the blood and empties the four chambers in a regularly repeated order. Apart from diseased conditions, the blood is driven in

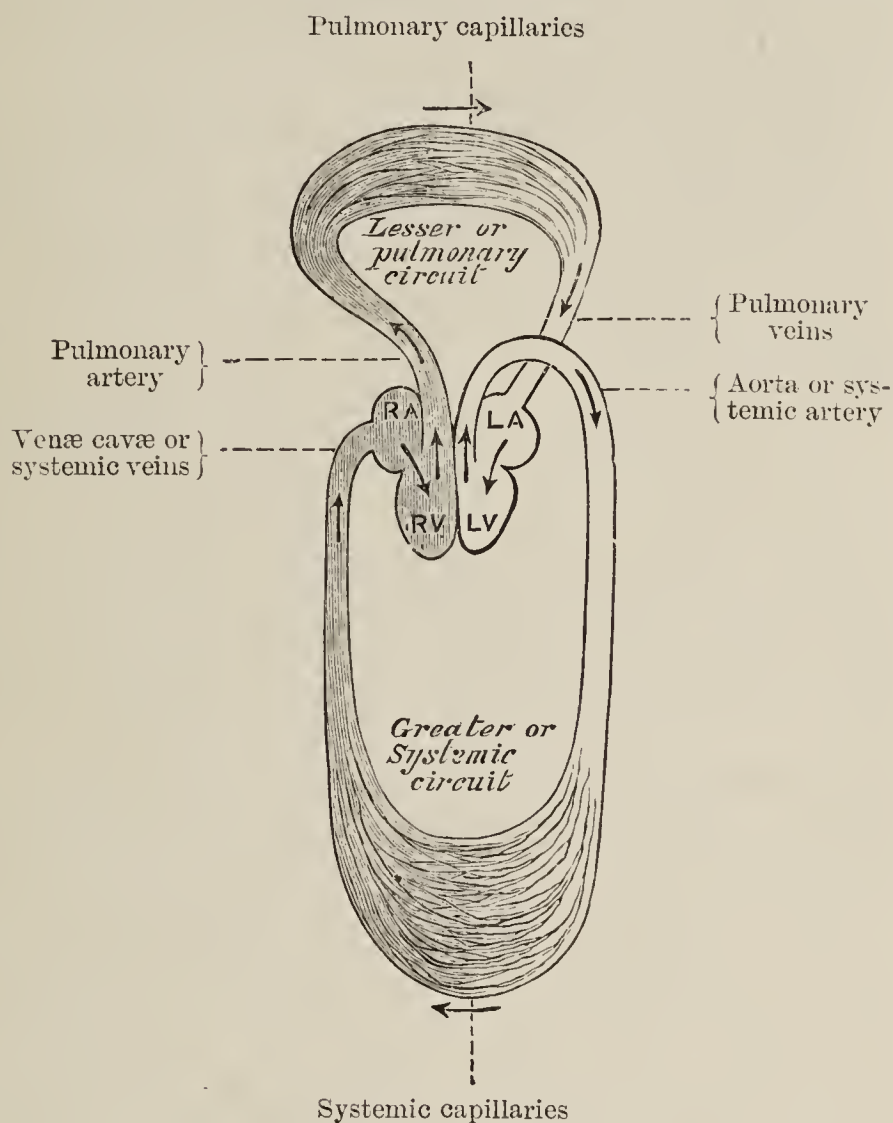


FIG. 13.—DIAGRAM OF THE CIRCULATION.



only one direction, namely, from auricles to ventricles, and from ventricles to arteries, and is prevented from flowing back in the opposite direction by valves. The *aortic semilunar valves* and the *pulmonary semilunar valves*, at the origin of the aorta and pulmonary artery, prevent reflux of blood from arteries to ventricles, while they offer no obstruction to the passage of blood from ventricles to arteries. The *mitral valves* between left auricle and left ventricle, the *tricuspid valves* between right auricle and right ventricle, prevent reflux of blood from ventricles to auricles, while they offer no obstruction to the passage of blood from auricles to ventricles.

The human heart contracts with regularity at a rate of about seventy per minute, and its parts act in regular sequence, each contraction or active period being succeeded by a passive period during which the previously contracted chamber is relaxed and becomes refilled with blood. This regular sequence of action and rest is termed the *rhythm*, and is of the following character: a short contraction of the auricles, immediately followed by a somewhat more prolonged contraction of the ventricles, the two auricles contracting together and the two ventricles contracting together. Each cardiac revolution or cycle consists of: (1) a short sharp contraction of the two auricles; (2) a longer contraction of the two ventricles; (3) a period of rest during which the heart is becoming refilled. The contraction of any part of the heart is called its *systole*; the opposite state, *i.e.* relaxation, is called its *diastole*; thus auricular systole signifies contraction of auricles, auricular diastole their relaxation; ventricular systole signifies contraction of ventricles, ventricular diastole their relaxation; and when the terms 'systole' and 'diastole' are used alone, or when 'cardiac systole' or 'cardiac diastole' are spoken of, it is the state of the ventricles to which reference is made.

Successive beats of the ventricles drive successive charges of blood into the arteries; and produce a *pressure* of blood within them: if an artery be opened the blood spurts from it in jets; if an artery be weakened by disease the pressure of the blood may burst it. There is no opportunity for blood to escape backwards from the arteries to the ventricles during diastole, because the semilunar valves are closed; it can only move onwards from aorta to large trunks, thence to branches, and finally to the small twigs called arterioles, and to the minute channels called capillaries. It is through these channels that the distended



arterial system is relieved, and it is owing to the *resistance* which these narrow channels offer that the arterial system is distended at all, and a pressure kept up within it. The greater or systemic circuit offers a greater resistance than does the lesser or pulmonary circuit; more force is required to drive the blood through the former than through the latter; parts concerned in driving and carrying blood through the system are more bulky and more robust than parts concerned in conveying blood through the lungs; thus the left ventricle has walls three times as thick as the right ventricle; the aorta and the systemic arteries are much thicker than the pulmonary artery and its branches; the aortic are much coarser and stronger than the pulmonary valves; the mitral curtains and their cords coarser and stronger than the tricuspid. The same difference, but much less pronounced, marks the auricles; the left is somewhat stronger than the right, but not much so; both are thin in comparison with the thinner of the two ventricles, the reason being that the auricles do not have to contract against resistance.

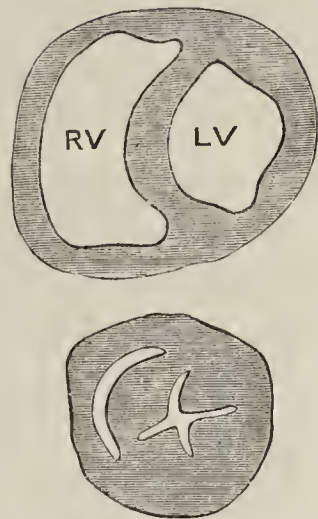


FIG. 14. — TRANSVERSE SECTION THROUGH THE MIDDLE OF THE VENTRICLES OF A DOG'S HEART IN DIASTOLE AND IN SYSTOLE (After Ludwig).

Beyond the resistance that is offered by the minute arteries and capillaries, *i.e.* in the veins, the pressure is very low—barely sufficient to send the blood onwards towards the heart—muscular movements, and in particular the movements of respiration, helping to this result; many veins have valves which, while they allow blood to move onwards towards the heart, do not allow it to move backwards towards the capillaries, so that if a vein full of blood is compressed by contracting muscles, its contents are squeezed towards the heart. In the lower extremities this disposition is particularly prominent, and has, as its effect, that the column of venous blood with each muscular movement is pressed upward from valve to valve, as up a ladder.

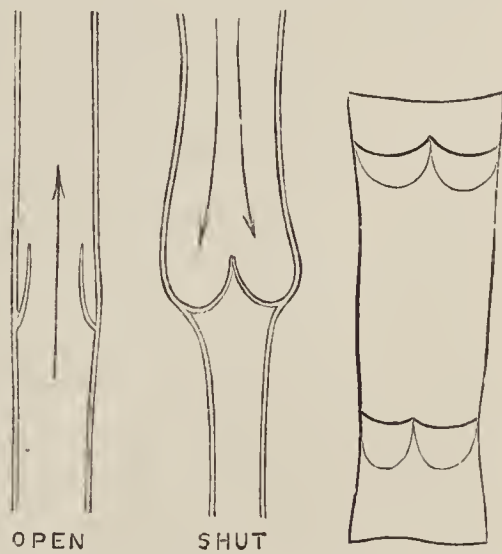


FIG. 15.—VENOUS VALVES. TO RIGHT OF FIG. A VEIN LAID OPEN TO EXPOSE THE POCKET-SHAPED VALVES.

The heart's movements are involuntary, and will even con-

tinue for a short time after the heart has been removed from the body; they may be accelerated or retarded through nervous channels; the escape of blood into the capillaries may be facilitated or obstructed by the relaxation or contraction of the muscular fibres which constitute the chief bulk of the walls of the minute arteries—such relaxations and contractions are for the most part effected through nervous channels. Thus the heart and vessels are not set for life to do uniform duty, but can be played upon and modified in action according to the requirements of special circumstances.

**The heart.**—*Physiological anatomy.*—Our knowledge of the physiology of the heart is based upon the study of the living organ in all classes of animals, but more especially in mammalia, such as the dog, cat, rabbit, horse, and man himself, and in cold-blooded animals, such as the frog, toad, and tortoise.

In mammalia the organ is constructed upon the plan above described—viz. it consists of two auricles or ante-chambers leading into two ventricles. It is essentially composed of *muscle* of a peculiar character, and is externally surrounded by a fibrous bag, the *pericardium*, internally lined by a delicate membrane, the *endocardium*. The *pericardium* is disposed in two layers, a visceral and a parietal layer, and these two layers are closely applied to each other, lined by a smooth membrane of endothelium, and barely moistened by serous fluid, so that with the movements of the organ the two surfaces glide upon each other without friction. This is in the healthy state; but when, in consequence of inflammation, the pericardium has become rough, the movements of the heart cause a friction between the two roughened surfaces, which may be heard or even felt, and the natural moisture may become exaggerated into a large amount of serous fluid.

The *endocardium* in health is a perfectly smooth membrane, lining the cavities of the heart and covering all its valves. As one of the chief consequences of its inflammation, it may become rough and irregular, so that the valves no longer act properly, and neither allow the blood to pass smoothly in the right direction, nor efficiently prevent its return in the wrong direction. Under such circumstances the natural sounds are obscured or replaced by noises or ‘bruits.’

The *muscle* of the heart, as examined under the microscope, differs from all other muscle, and occupies an intermediate position between involuntary or unstriped muscle and voluntary



or striped muscle. It is composed of short, four-sided cells with short, thick branches, which join with the branches of other cells, thus forming a closely interlacing network; the cells are striated across their length, they possess nuclei deeply embedded in their substance, and they are devoid of any surrounding membrane (sarcolemma).

The *bulk* of muscle bears an evident relation to the amount of *work* that falls to the share of the several chambers of the heart. Thus, as already mentioned, the left ventricle is thicker than the right, and the auricles are thin in comparison with the ventricles. A similar relation between work and bulk holds good in diseased states, when, from any cause, the circulation is embarrassed and effected with unusual difficulty. A most common cause of excessive enlargement of the heart is the imperfect action of a valve, which either obstructs the free exit of blood from a cavity, or allows blood to return in consequence of imperfect closure; such an imperfection is equivalent to an addition of work, and the consequence of additional work is increased bulk of muscle. An excessive growth (*hypertrophy*) takes place gradually in response to the necessity, and the imperfection may thus be adequately compensated for a long period.

The *nervous supply* of the heart is derived from the medulla oblongata and spinal cord by two channels—the *vagus* and the *sympathetic*. Both kinds of nerve-fibres, medullated as well as non-medullated, are contained in these nerves, and numerous ganglion-cells, in some places collected into ganglia of considerable size, are situated on their course.

The hearts of cold-blooded animals, that of the frog especially, owe their physiological importance to the fact that they are easily studied. The heart of a cold-blooded animal—frog, fish, tortoise, crocodile, &c.—will continue to beat for hours after it has been cut out of the body, whereas the time during which the heart of a mammal will beat under similar circumstances is to be counted in minutes only. It is necessary to notice the points in which the hearts of cold-blooded animals differ anatomically from those of mammals, and it is advisable in the appreciation of physiological results to distinguish between such as refer only to cold-blooded animals, and others that refer only to mammals.

Thus the experiment of the ‘Stannius ligature’ refers to the frog’s or toad’s heart; the action of the depressor nerve is experimentally known only in the case of the rabbit’s or cat’s heart; observations of intra-cardiac pressure have been mostly



made on the dog's or horse's heart; the characters of the heart's sounds and of the pulse are studied in detail only in the case of man.

*Structure of the frog's heart.*—The *frog's heart* is composed of a single ventricle and two auricles. The right auricle receives the venous blood of the whole body; the left auricle receives the blood of the lungs; the venous blood of the body is carried to the sinus venosus through the three venæ cavæ, and thence into the right auricle. Both auricles open into the ventricle, which thus receives mixed blood. From the ventricle the blood is driven into the bulbus arteriosus, and thence into the system, the lungs, and the skin. The mixed blood from the ventricle loses oxygen in the system, gains oxygen in the lungs and skin. The muscular fibre of the frog's heart differs from that of the

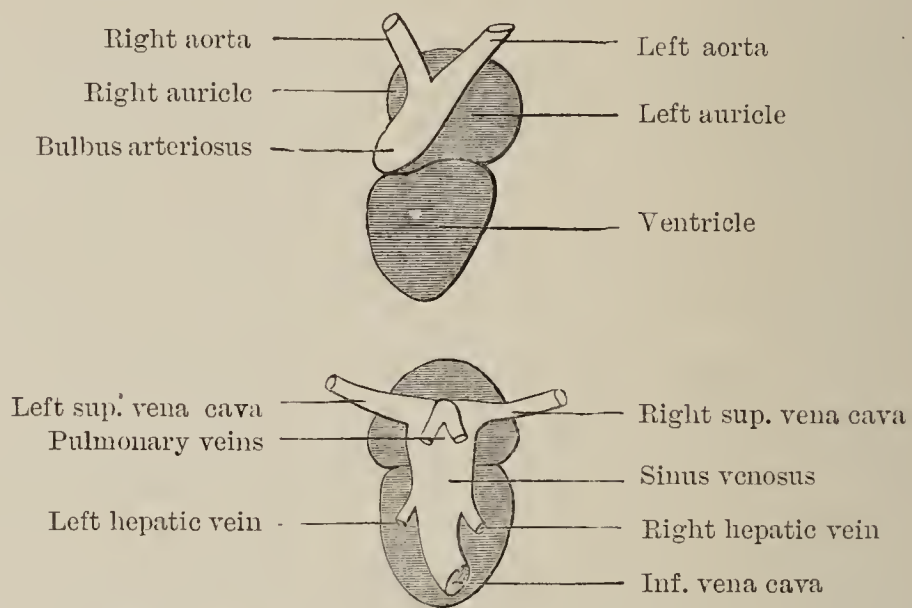


FIG. 16.—ANTERIOR AND POSTERIOR VIEWS OF FROG'S HEART. (After Cyon.)

adult mammalian heart; it is of a more embryonic character, composed of long spindle-shaped cells, which are nucleated and dimly cross-striated, closely resembling the fibre of the embryonic mammalian heart. The cavity of the ventricle is intersected in all directions by bundles of muscle, giving the mass a spongy structure; this is well seen when an inflated and dried ventricle is cut across. Nerve-fibres, medullated and non-medullated, the former derived from the vagus, the latter from the sympathetic, are abundant in the auricles and in the upper third of the ventricle. Nerve-cells, scattered and massed together as ganglia, are abundant in the same situations; they are probably in connection with medullated or vagus fibres; their principal masses are in the sinus venosus (Remak's ganglion), in the auricular septum (von Bezold's ganglion), and at the base of the ventricle (Bidder's

ganglion). Nerve-fibres, but no nerve-cells, have been detected in the lower two-thirds of the ventricle.

The heart of man is habitually examined in hospital wards by various instruments applied to the chest-wall where the heart rests against it. The character of beat, as regards *force* and *frequency*, is judged of by mere inspection, or by the hand laid upon the chest-wall. The size of the heart is ascertained roughly by *percussion*. The sound which percussion of the chest-wall brings out is different according to the nature of what lies behind it: where the heart lies behind the chest-wall the percussion-sound is far duller and less resonant than where the lung lies, and an area of 'dulness' may thus be mapped out, the extent of which gives some notion of the size and position of the heart.

But the most important indications are those furnished by the *sounds* of the heart, which reveal the condition of its valves; if the ear is applied to the front of the chest the sounds of the heart will be plainly heard—two sounds with a rhythm resembling that of the footsteps of a man who walks lame. The *first sound* is muffled and prolonged in comparison with the second sound, which is short and sharp. Their character is imitated by the syllables *lub-dup*, but can only be properly realised by applying the ear to a person's chest. To simply hear the heart's sounds this is sufficient: to localise on the chest-wall the spots where the sounds are best heard, a 'stethoscope' must be used. The sound due to closure of the mitral valve is loudest when the stethoscope is placed over the 'apex-beat'; that due to closure of the aortic valves is loudest at the second right costal cartilage. It is then easy to recognise that the first sound is simultaneous with the impulse of the heart; it occurs with the ventricular systole, and is therefore called *systolic*; thus the impulse gives an easy means of recognising the first from the second sound, if the distinction has not already been made by means of their rhythm and character. The second sound occurs at the end of the ventricular systole, or, in other words, at the beginning of the ventricular diastole, and it is therefore called *diastolic*. As to the cause or causes of the sounds, these have been ascertained by experiments on animals and by observations on man. The first, or systolic sound, is due to two causes combined: (a) the sudden closure of the auriculo-ventricular valves; (b) the noise of muscular contraction. The second, or diastolic sound, is caused by the sudden closure of the semilunar valves. The latter statement has been proved as



follows : curved needles have been introduced through the carotid artery into the aortic orifice, and the aortic semilunar valves have been hooked back so that they could not close properly ; the second sound has then lost its sharp character and become replaced by a rushing noise. On withdrawal of the needle the sharp slapping second sound has returned ; therefore the second or diastolic sound is due to the sudden closure of the semilunar valves. If the valves have become permanently injured, so as to be incompetent to close the orifice when it should be closed—*i.e.* when diastole begins—the character of the second sound is permanently altered, the sharp slap is replaced by a diastolic noise, which is caused by the blood as it escapes back into the ventricle. On man such diastolic noises or murmurs are not unfrequently heard instead of a natural second sound, in consequence of an incompetent state of the aortic valves, and the fact is frequently verified after death, thus affording confirmation of the statement that the second sound is due to the sudden closure of the semilunar valves. That the first, or systolic, sound is caused by muscular contraction and by valve-closure combined is proved by two experiments : (1) The closure of the mitral valve has been prevented by a wire ring introduced into the auriculo-ventricular orifice—the first sound was thereby much altered ; or the chordæ tendineæ have been divided by a curved knife, with a similar result to the first sound. (2) A systolic sound can still be heard in the excised and empty hearts of dogs. There is here no possibility of valve-tension, and the noise made by the contracting ventricle must therefore be admitted to be a factor in the production of the first sound. (3) On man a natural first sound is not unfrequently modified or replaced by a systolic murmur ; one of the causes of this, as has been verified after death, is an incompetence of the mitral valve, which does not close properly, but allows blood to escape backwards into the left auricle during the ventricular systole.

The heart's *impulse* is usually most easily felt in the fifth intercostal space, about one inch below the nipple and one and a half inch nearer the sternum. The situation of the impulse is, however, by no means invariable ; it may be more marked in the fourth intercostal space, or further towards the axilla than the nipple ; it is altered with the movements of respiration and with the position of the body, and it is modified both in situation and extent by variation in the size of the heart. This spot, where the impulse is greatest, is often termed the 'apex-beat'—



a misleading expression, for the true apex of the heart lies much deeper ; a needle plunged through the spot of maximum impulse would pierce the left ventricle at the junction of the middle with the lower third. The cardiac impulse is not produced by the heart's tilting itself so as to strike the chest-wall ; it is due to the sudden hardening and tension of the contracting ventricles, and the spot of impulse, or so-called 'apex-beat,' is simply the spot where the convex ventricular mass comes in contact with the chest-wall.

The impulse of the heart may further be studied by means of the *cardiograph*. This is an instrument by means of which the movement produced by the impulse is magnified and recorded as a curve on a

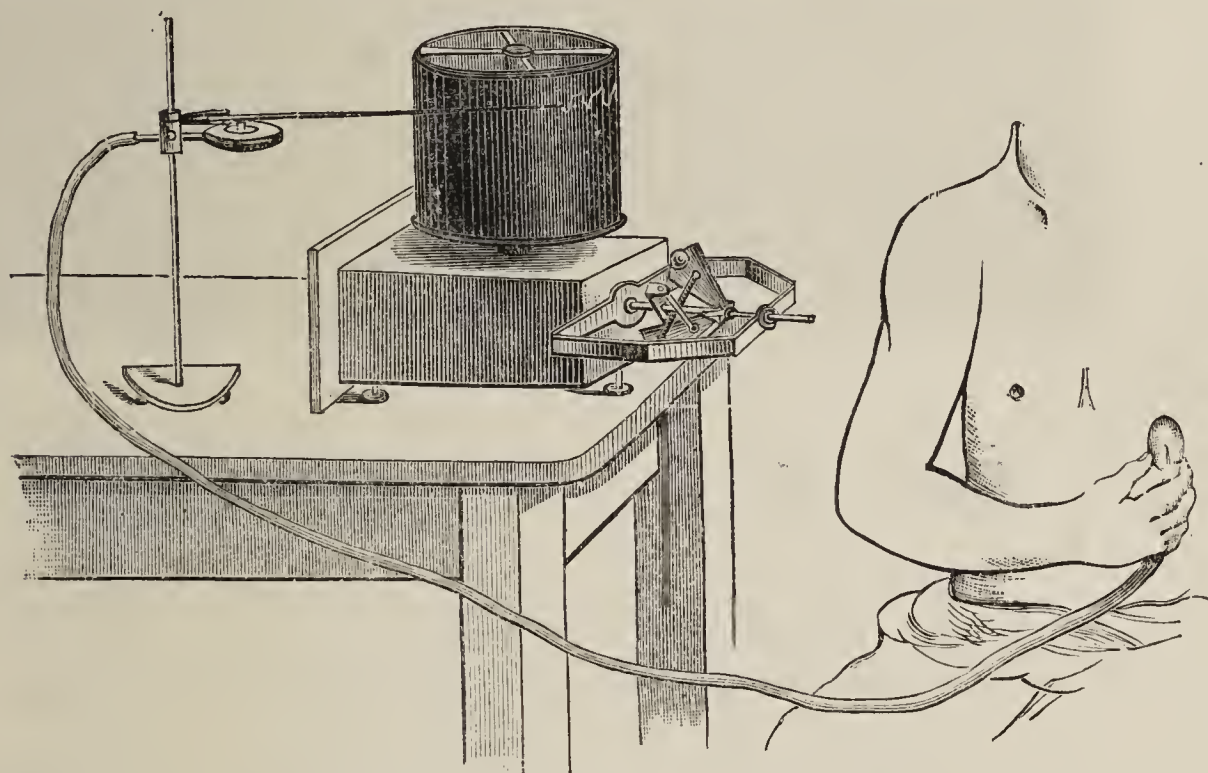


FIG. 17.—THE CARDIOGRAPH AS APPLIED TO MAN.

travelling surface. A cardiograph consists of the following parts : (1) an exploring tambour, which is applied to the spot where the impulse is best felt ; this tambour is joined by a tube with (2) a recording tambour which carries a lever. These tambours are like small kettle-drums, with drumheads made of a very elastic membrane. Both tambours and connecting tube are full of air, and any pressure, such as that produced by the heart's impulse, exerted upon the membrane of one tambour, is transmitted through the connecting tube to the other tambour causing bulgings of its membrane, which are magnified by the lever. The lever marks its movements against (3) a travelling surface, a usual form of which is a cylinder covered with smooth paper and blackened by a smoky flame ; the point of the lever rubs lightly against this blackened surface and makes a white line, or 'tracing.' The rate

at which the cylinder revolves varies according to the special requirements of observation ; it is determined by means of a time-marker, or 'chronograph,' which is essentially a lever also touching the smoked paper, and moving once per second, so as to mark seconds ; or 10 times per second, so as to mark tenths of a second ; or 100 times per second, so as to mark hundredths of a second, &c. (p. 316).

By means of the cardiograph we may, with due precaution, very accurately determine the time occupied by the systole of the auricles and of the ventricles, and by their diastole ; a *typical* apex cardiogram corresponds in duration with that of the intra-ventricular pressure (fig. 23). The following figure gives an example of a heart-tracing thus obtained, the lower line being

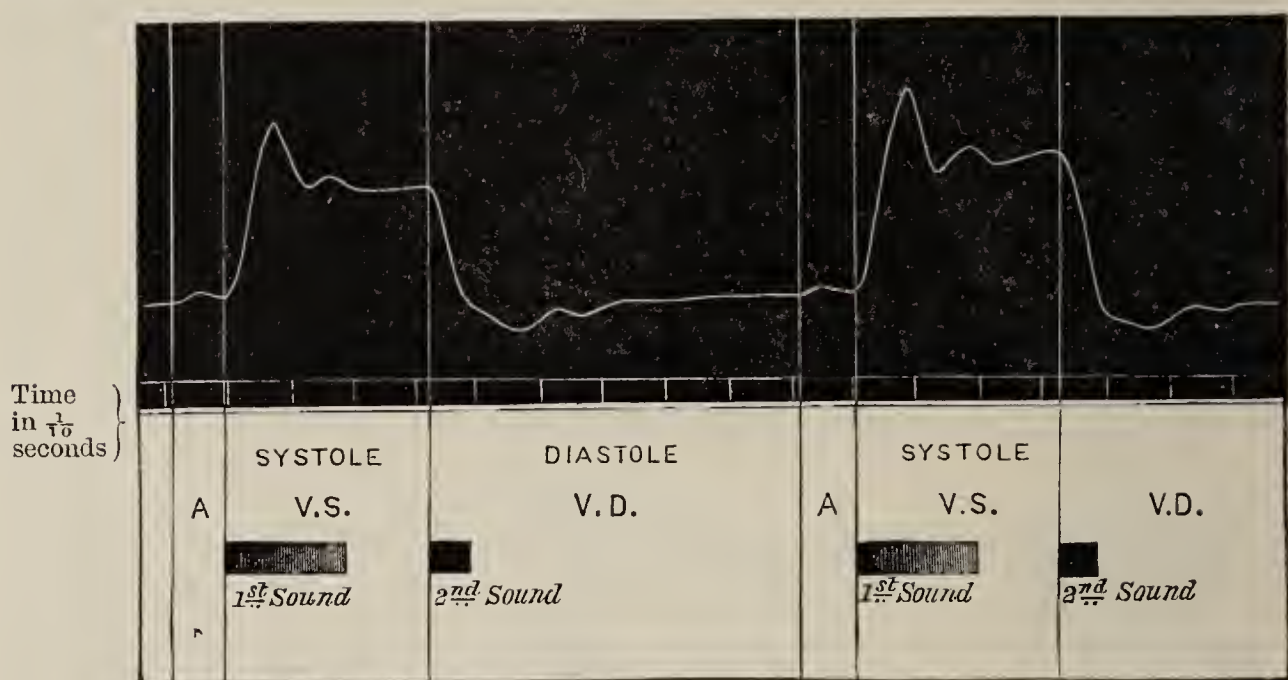


FIG. 18.—A TRACING FROM THE 'APEX-BEAT' OF MAN.

Relation of sounds to contraction indicated below the tracing by black marks.

the time marked in tenths of a second. The auricular systole is seen as a small elevation immediately preceding the larger and longer elevation caused by the ventricular systole, the beginning and end of which are shown by the points at which the lever begins to rise and begins to fall.

If the movements of the recording lever be carefully watched while the heart's sounds are listened to, it will be apparent that the first sound is heard as the lever rises, the second sound as it falls—*i.e.* the first sound is systolic, the second sound is diastolic. Simultaneous records of the heart's beat and of the heart's sounds (of the latter by means too complicated for brief description) give further and perhaps more indisputable evidence that the sounds take place precisely as stated (and diagrammatically represented in fig. 18). Assuming that the heart is beating sixty times per



minute, so that each cardiac event or cycle lasts one second, the systole and diastole of the auricles and ventricles will occupy the following fractions of a second :—

Auricular systole	.	.	.	.	.	one-tenth
Auricular diastole	.	.	.	.	.	nine-tenths
Ventricular systole	.	.	.	.	.	four-tenths
Ventricular diastole	.	.	.	.	.	six-tenths

An increased number of beats per minute naturally implies diminished length of each diastolic pause, also—but in less degree—diminished length of each systole. Apart from alterations of frequency, the duration of systole is not appreciably affected by variations of arterial pressure. With each increase of ten beats per minute, the diastole decreases by about  $\frac{1.0}{100}$  of a second in duration, the systole by about  $\frac{2}{100}$ . Consequently the total systolic or working time of the heart, as compared with its total diastolic or resting time, is greater at a high than at a low pulse-frequency. Normally the human heart is at work for about nine hours of the twenty-four, at rest during about fifteen hours—points that are exhibited in fig. 19 and in the accompanying table.

The foundation of all precise acquaintance with the mechanism of cardiac contraction—abnormal as well as normal—is summarised in the following propositions, each of which must be realised and mentally ‘seen’ before the details of individual cases and experiments can be understood. (1) *During systole the ventricles contract and empty themselves into the pulmonary artery and aorta; the semilunar valves of these vessels are open; the auriculo-ventricular valves are shut; the first sound is produced.* (2) *During diastole the ventricles are relaxed; the semilunar valves of the aorta and pulmonary artery are closed; the auriculo-ventricular valves are open, and the blood is flowing from auricles to ventricles; just at the end of the diastole the auricles contract. The second sound is produced at the beginning of diastole.*

These are the main facts, around which all others attach themselves as accessory and subordinate. Among these accessory facts are: the alterations of volume in systole and in diastole, the rotation of the contracting and relaxing ventricle, the mode of commencement of the auricular contraction, the manner in which the muscoli papillares act upon the intra-cardiac valves and secure their perfect closure. These points may be briefly disposed of. The volume of the heart is diminished



during systole, the ventricular mass in contracting becomes shorter as well as narrower, although on superficial examination the narrowing is such as to give an appearance of elongation.

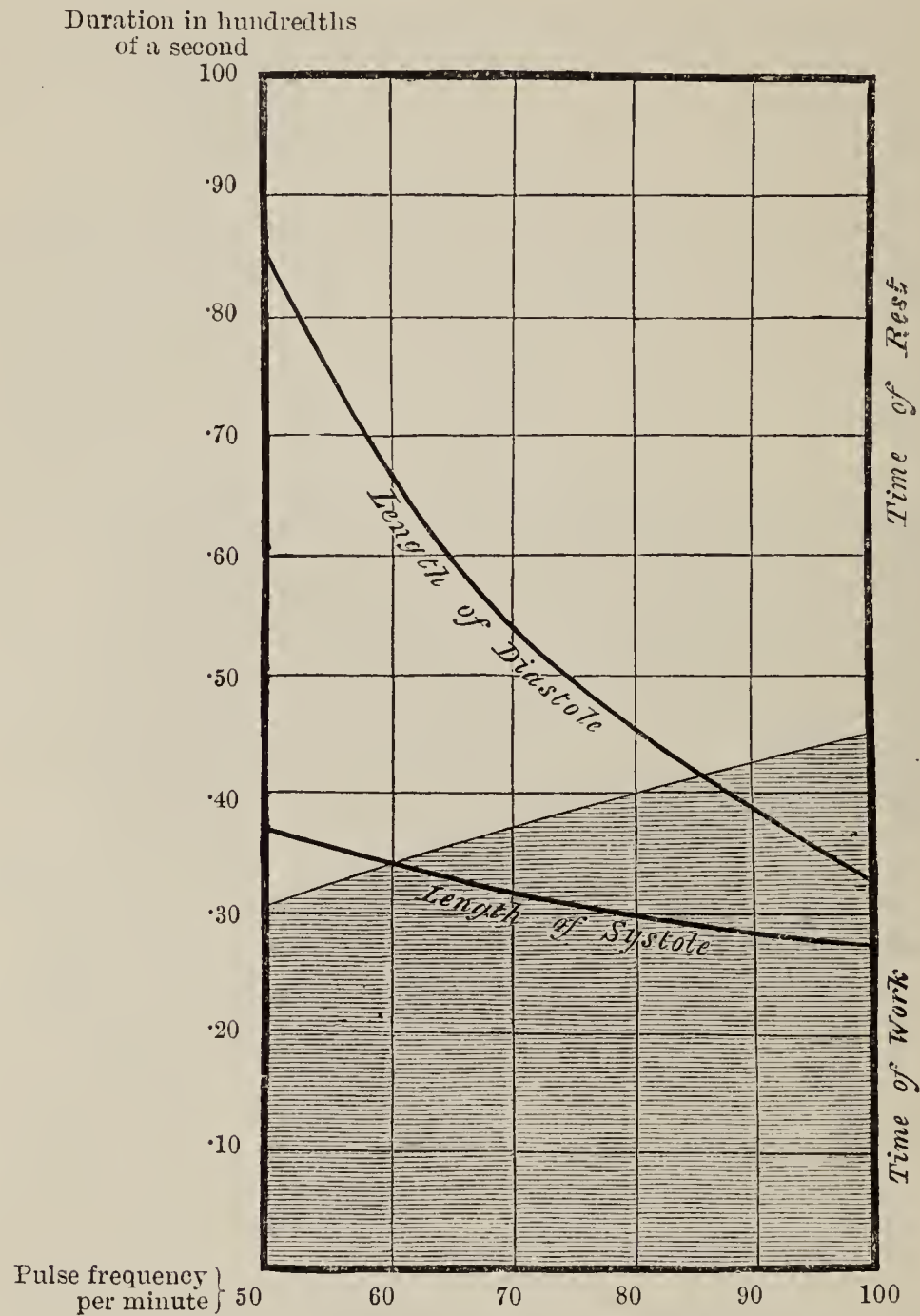


FIG. 19.—DURATION OF SYSTOLE AND OF DIASTOLE WITH DIFFERENT PULSE-FREQUENCIES. (W.)

Pulse-frequencies per minute are indicated along the abscissa. Durations of systole and of diastole are given in hundredths of a second. Their respective curves show that the systole shortens by about  $\frac{2}{100}$  second for each increase of 10 beats per minute, and that the diastole shortens by about  $\frac{10}{100}$  second.

The shaded and unshaded portions represent respective time of work and time of rest at various pulse-frequencies.

Pulse-frequency per minute	Duration of sys- tole in hundredths of a second	Duration of dia- stole in hundredths of a second	Ratio of systole to cardiac cycle	Hours of work per diem
50	37	83	.31	7.5
60	34	66	.34	8.2
70	32	54	.37	8.9
80	30	45	.40	9.6
90	28	38	.42	10.2
100	27	33	.45	10.8

If the auricles be closely watched when the heart is beating slowly, it will be noticed, in the left auricle especially, that the contraction starts from the venous orifices, and that it is in fact ushered in by contraction of the large veins; the auricles are anatomically as well as physiologically continuous with the large

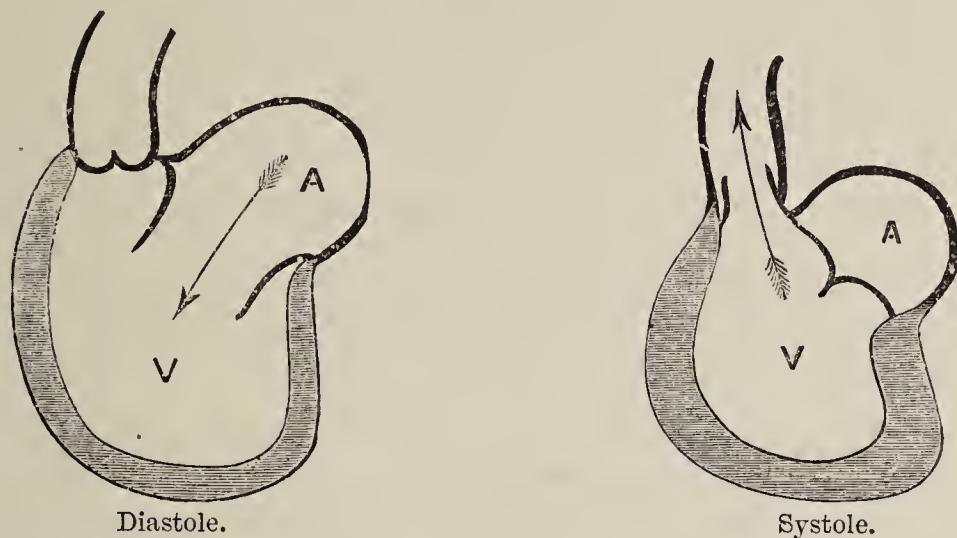


FIG. 20.—DIAGRAM TO ILLUSTRATE THE STATE OF THE CARDIAC VALVES AND OF THE BLOOD-CURRENT DURING THE DIASTOLE AND SYSTOLE OF THE VENTRICLES.

veins which open into them. The points so far considered may be verified by simple inspection of the moribund and therefore sluggishly-beating heart of any recently killed mammal; the exact action of the muscoli papillares under normal conditions has been until quite recently a subject of argument rather than of direct observation; from their attachment to the inner surface of the ventricle near the apex, and by the chordæ tendineæ to the free edges of the curtain-like cusps of the mitral and tricuspid valves, it is obvious that they must prevent these cusps from being floated back into the auricles when the ventricles begin to contract; by their contraction they must compensate the shortening of the ventricle and thus keep the valves tight. According to Roy and Adami the papillary muscles begin to contract a little later, and cease to contract a little sooner than the rest of the ventricle.

**Blood-pressure.**—At each stroke of the ventricles blood is driven into the arteries; its passage into the veins is impeded by the friction of the mass of blood against the walls of the capillaries, and by the narrowness of the minute arteries leading to them. The resistance thus offered is spoken of as the ‘peripheral resistance.’

The *arteries* are distensible and elastic—that is to say, they can be distended by internal pressure, and they can shrink again when the internal pressure ceases. Elasticity should be carefully distinguished from distensibility or extensibility. A band



of india-rubber lying loose on the table is extensible—it may be taken up and stretched, but it is not elastic until it is stretched ; while held stretched between two hands it is elastic—that is to say, it is pulling the two hands together, and will shorten if let go. The arteries can be stretched by internal pressure, *i.e.* they are distensible ; further, the arteries are in the living body kept stretched by the blood driven into them, *i.e.* they are elastic ; and their elasticity is constantly exerted in pressing upon the blood. The *blood-pressure* is thus the resultant of two factors—(1) the heart's force, (2) the peripheral resistance. The elasticity of the arteries may be named as a third factor, but it must be borne in mind that this recoil force is not an independent factor ; it is derived from the heart's force. Arterial elasticity supplies no fresh force, but simply acts as the reservoir of the heart's force, just as the distended air-bag of a piper acts as the reservoir of his expiratory efforts ; its effect as regards the circulation is to convert pulsating, intermittent force into continuous force ; each beat of the heart is in great part spent in keeping the arteries distended, bringing into play their elasticity, which is constantly compressing the contained blood, thus keeping up a constant flow in the intervals between the beats. A part of each beat is, however, not thus spent in keeping up elasticity and pressure, but goes directly to increase pressure and onward flow ; it is this portion which is manifested as the pulse, and in a corresponding acceleration of the velocity of the blood-current. The arterial blood-pressure will obviously vary with variations of its factors—*i.e.* pressure will be greater with greater heart's force or with greater peripheral resistance, and *vice versa*.

*The Manometer.*—Blood-pressure is measured on animals by means of the *manometer*, of which there are various forms ; the simplest and most convenient is the mercurial manometer. A bent glass tube of suitable length, half filled with mercury, is connected with the interior of an artery, say the carotid, by means of a tube and cannula filled with sodium carbonate solution. The blood presses through the cannula and tube towards the manometer, and causes the mercury to fall in one limb and rise in the other. The difference of level at which the mercury stands in the two limbs gives the pressure of mercury equal to the pressure of blood in the case examined. If the column of mercury be closely watched, or if a float carrying a pen be arranged so as to record the level of the column upon a revolving cylinder, it will be noticed that the height of the column is variable—irregularly variable, *i.e.* rising or falling more or less without obvious cause or in consequence of muscular efforts ; and regularly variable, rising and falling



with each beat of the heart and with each act of respiration. The complete apparatus for taking blood-pressure records is commonly termed a *kymograph* (fig. 21).

Arterial blood-pressure cannot be studied upon man by the method above described; all direct evidence on the subject is derived from experiments on animals, such as the dog, rabbit, and horse, and from the facts thus obtained we judge of what must take place in the human arterial system. The mean blood-pressure in the carotid artery of a rabbit is equal to about 8 to 10 cm. of mercury, of a dog or horse 12 to 15 to 20 cm.; in the human carotid artery the pressure is supposed to have about the same value, viz. about 15 to 20 cm. (=6 to 8 inches).

Attempts are, indeed, made to estimate arterial pressure clinically by means of the sphygmograph and other instruments, such as the plethysmograph and the sphygmomanometer, but it is difficult to get exact results by any instrument, and the pulse skilfully felt is the readiest means of estimating arterial pressure on man. By feeling the pulse, we may ascertain that blood-pressure is unusually high, or unusually low, or not far wrong; we say, accordingly, that tension is high or low or normal, and we speak of a pulse as of high tension, or of low tension, or of normal tension.<sup>1</sup>

We have recognised as obvious that, *cæteris paribus*, arterial pressure should be greater with greater heart's force, less with less heart's force, greater with greater peripheral resistance, less with less peripheral resistance. All these statements may be illustrated by experiments and observations. Violent exertion causes the heart to beat more frequently (and possibly each beat may be stronger, though this is not necessarily so), but in any case the total force exerted by the heart during a given time—say a minute—is greater than usual, and the blood-pressure is temporarily increased. A sudden shock, or a strong emotion, or experimental stimulation of the vagus nerve, weakens or arrests the action of the heart, and the blood-pressure is temporarily diminished. If the vasomotor centre is stimulated, the muscular

<sup>1</sup> The word 'tension' is more commonly used in clinical medicine than the word 'pressure.' Arterial tension signifies the elastic force exerted by a distended artery; this elastic force is equal to the distending force of blood-pressure. The term 'arterial tension,' 'arterial pressure' may therefore be substituted each for the other. The term 'blood-pressure,' used without qualification, is understood to denote arterial blood-pressure.

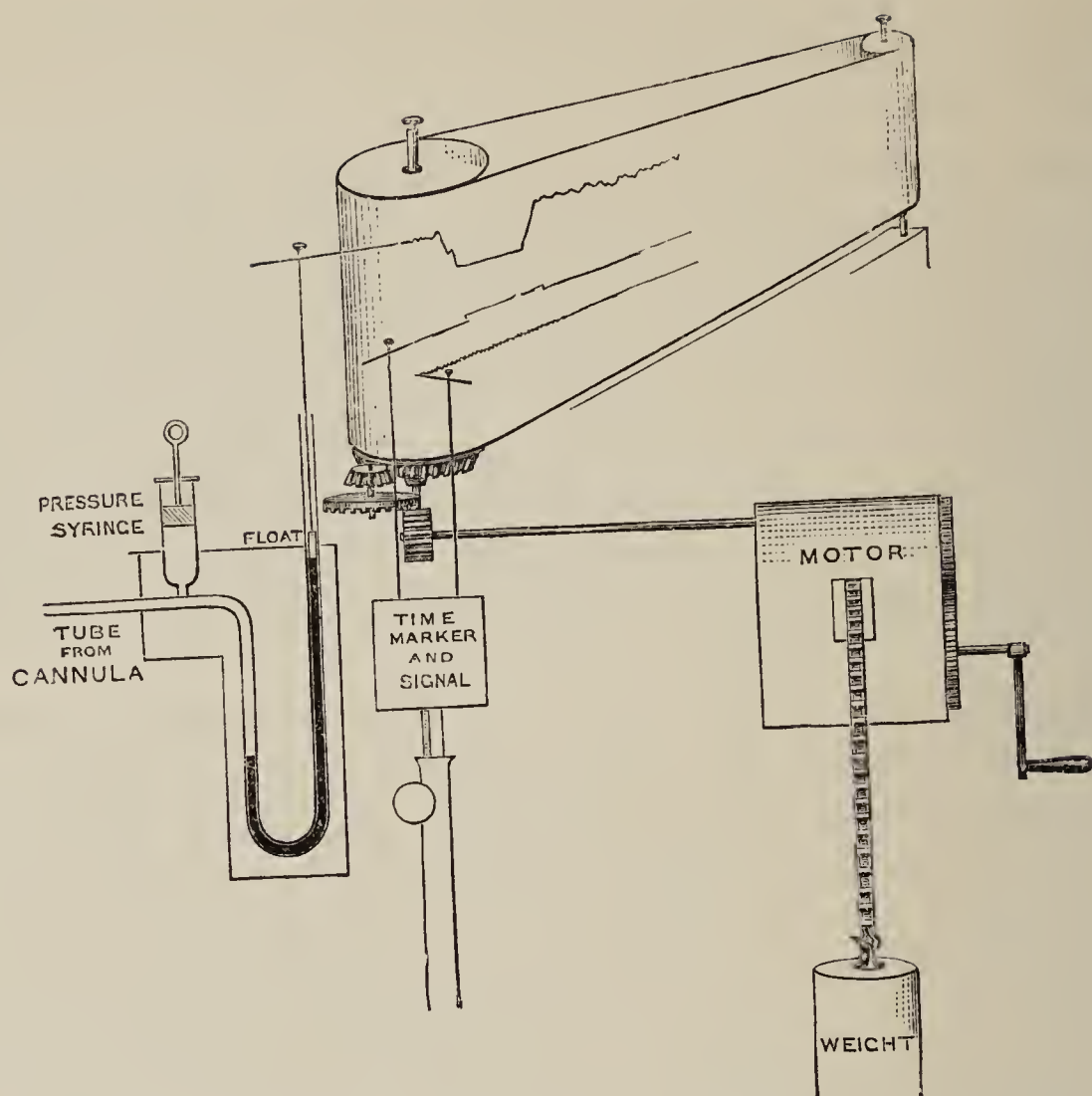


FIG. 21.—DIAGRAM SKETCH OF THE MERCURIAL KYMOGRAPH. (Hering's Model.)  
(κῦμα, wave; γράφω, to write.)

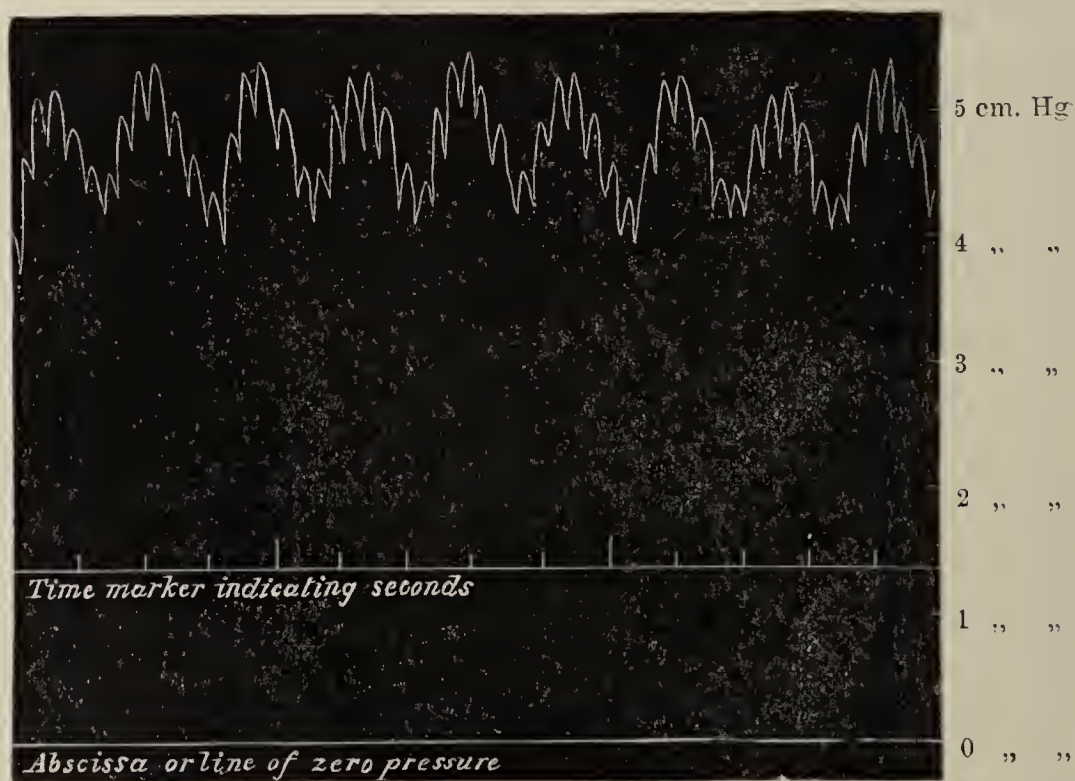


FIG. 22.—PORTION OF A BLOOD-PRESSURE TRACING FROM THE CAROTID ARTERY OF A RABBIT.

The small undulations are due to the heart-beat; the larger undulations upon which they are superposed are due to the movements of respiration. The height above the zero line is one-half the blood-pressure expressed in terms of mercurial pressure.



arterioles contract and narrow the outlet from the arterial system, so that the peripheral resistance is increased; blood-pressure is raised. If the vasomotor centre is destroyed, or its action is depressed by shock, the muscular arterioles relax and widen the arterial outlet; blood-pressure falls.

*Blood-flow.*—All these are extreme cases, which actually do occur in the living body; but we have to recognise that the several alterations, whether of the heart or of the arteries, naturally limit and neutralise each other, so that a marked change of pressure, as above described, can only occur in consequence of an excessive alteration of one of two factors. The co-operative variations of the two factors will be better understood by taking into account how the *rapidity of the blood-flow* alters with the alterations of the heart or of the vessels. For the sake of brevity, these alterations may be cast into tabular form by the side of the alterations of blood-pressure which we are discussing. The table includes all the possible variations which may theoretically take place, the consequences in each case being denoted by the signs + or —, to signify increase or diminution.

No.	Heart	Arterioles	Blood-pressure		Blood-flow	
1	{ Force constant.....	Resistance increased	+		—	
2		Resistance diminished	—		+	
3	{ Force increased .....	Resistance constant	+		+	
4		Resistance constant	—		—	
5	{ Force increased .....	Resistance diminished	+	—	+	+
6		Resistance increased	—	+	—	—
7	{ Force increased .....	Resistance increased	+	+	+	—
8		Resistance diminished	—	—	—	+

The first four cases need not detain us long. If—supposing the heart's force to remain constant—the arterioles contract and increase the peripheral resistance, it is obvious that pressure will be raised, and that through the constricted outlet less blood will flow (1); *vice versa*, if the arterioles dilate, widening the outlet and diminishing the resistance, it is clear that pressure will fall and more blood will flow through the vessels (2). If—supposing the peripheral arterioles to remain unaltered—the heart beats more strongly or less strongly, it is obvious that with more force there will be increased pressure and increased flow of blood (3), with less force there will be diminished pressure and diminished flow (4).



Turning to the next four cases, in which both factors are altered, it is evident that in (5) increased heart's force will give more pressure, while diminished peripheral resistance will give less pressure; these alterations neutralise each other, and pressure remains unaltered, or, if either should exceed the other, the resultant alteration, being only their difference, is not great. But as regards blood-flow the result is otherwise; increased heart's force gives increased flow, diminished peripheral resistance also gives increased flow, and these two alterations united give a resultant alteration which is comparatively great. Similar reasoning applies to the combination (6), all data and results being reversed; thus, diminished force of the heart gives less pressure and less flow, increased resistance gives greater pressure and less flow; as our resultants we have little or no alteration of pressure, and a great diminution of the blood-flow. Thus in these two cases (5 and 6) we have a minimum alteration of blood-pressure with a maximum alteration of blood-flow. If now we consider the last two cases, we shall find how differently the results come out. In combination (7) the increased heart's force gives increased blood-pressure, and so does the increased peripheral resistance; the resultant increase of blood-pressure is therefore considerable, while as regards the blood-flow the first factor increases, the second diminishes it, and the resultant alteration is little or nothing. In combination (8) similar reasoning applies; both factors diminish the blood-pressure, but the first diminishes while the second increases the blood-flow. Thus in these two cases we have a maximum alteration of blood-pressure with a minimum alteration of blood-flow.

The body, as a whole, requires, according to circumstances, more or less blood; more or less blood-pressure is of itself useless, and accordingly we find that the temporary variations naturally occurring in the body are such as produce a *minimum alteration of blood-pressure with a maximum alteration of blood-flow* (viz. cases 5 and 6), whereas the converse variations (viz. 7 and 8), giving a maximum alteration of blood-pressure with a minimum alteration of blood-flow, are abnormal and do not occur in the healthy body. In diseased conditions they do, however, occur, and persist as enduring states: in renal diseases a strongly beating heart with constricted arterioles, and consequently high blood-pressure, form a very usual combination; in great prostration from any cause, a weak heart with relaxed

arterioles, and consequently a low blood-pressure, is the invariable rule. And it may be incidentally noticed how in these cases the two factors add themselves as regards blood-pressure, but oppose and compensate each other as regards blood-flow, so that the body is not starved of blood in the same proportion as the circulation becomes weaker (case 8) ; nor over supplied with blood in proportion to a heightened blood-pressure (case 7). Referring to the table, we see that in these combinations a small alteration of blood-flow accompanies a great alteration of blood-pressure. Thus analysed we can recognise the economy of the relationship between the state of the heart and of the vessels, and between blood-pressure and blood-supply. The essential event is blood-supply, to which blood-pressure is subservient. Normal fluctuations of tissue-activity elicit corresponding fluctuations of blood-supply with minimum alterations of pressure. Abnormal alterations of blood-pressure do not, until they are excessive, interfere with a normal and necessary rate of blood-supply.

It should, however, be added that the immediate effects of increased or diminished blood-pressure are always an increase or a diminution of blood-flow, although in the first case the vessels are constricted and in the second case dilated. This result, which could not have been foreseen on *a priori* grounds, has been established by direct measurement of the aortic blood-current. Destruction of the spinal bulb relaxes the arterioles, lowers blood-pressure, and reduces blood-flow ; excitation of the bulb constricts the arterioles, raises blood-pressure, and accelerates blood-flow.

*Local variations.*—The general requirements of the whole body vary, and are met, as above described, by variations of the general blood-pressure and blood-flow, variations of flow being more important than variations of pressure. We have further to consider *local variations* in response to the particular requirements of its several parts or organs. Two different parts or organs may at the same time require very different amounts of blood, the same part or organ requires at different times very different amounts of blood. Such local variations of requirement are met by local variations of the peripheral resistance, the arterioles of the part or organ contracting when less blood is required, relaxing when more blood is required. A gland during secretion, a muscle during contraction, the digestive viscera during digestion, the brain during mental exertion, require and



receive more blood than the same parts while at rest; their arterioles dilate, the blood-flow through them is increased, more food is received during more work. Such local changes may or may not affect the general circulation, according to their magnitude or rate of occurrence. If the changes occur in a sufficiently large district—as, for instance, in the area governed by the splanchnic nerves, *i.e.* the intestinal vessels—the sequence of effects will be as follows: a contraction occurs of the splanchnic area, rise of blood-pressure in that area, and throughout the system—less blood passes through the splanchnic area, more blood passes through the remainder of the system; or a dilatation occurs of the splanchnic area, blood-pressure falls in that area and elsewhere throughout the body—more blood passes through the splanchnic area, less blood passes through the system. This last series of effects is, indeed, precisely what takes place after every copious meal; blood is then diverted to the intestinal vessels, there is less blood than usual in the remainder of the system; a feeling of chilliness and a disinclination to exertion, mental or physical, are the tokens that much blood is engaged in visceral action and cannot be spared to other parts (Kant); or if the disinclination be overcome, if by forced exertion blood be called to the brain or to the limbs, then the visceral blood-flow is made insufficient, and digestion is disturbed.

*Addition and removal of blood.*—It might appear at first sight that the blood-pressure should be relatively high when the system contains more blood than usual, relatively low when it contains less blood than usual. Yet in the living body this does not necessarily hold good; a large quantity of fluid may be injected into the arterial system without causing any increase of blood-pressure, and an animal may be copiously bled without lowering the blood-pressure. It is not until the excess or deficiency is pushed to extremes—*i.e.* the total amount nearly doubled or halved—that any effect is produced, and then death usually follows. These facts are of importance, for they show that the arterial system, by contraction or by relaxation of its muscle, can accommodate itself to very different quantities of blood without alteration of pressure, and that we should not attempt to reduce a high blood-pressure by removing blood, or to raise a low blood-pressure by injecting blood or other fluid. The total quantity of fluid present in the vascular system is, moreover, kept very uniform by osmotic currents, which transfer water from lymph to blood after a loss of blood, or from blood



to lymph after an injection of fluid into the vascular system. In as short a period as half an hour after copious hæmorrhage, the dilution of the remaining blood is recognisable by a marked diminution of specific gravity; and the opposite effect—viz. recovery from dilution after an injection of fluid—is seen in the hardly less rapid restoration of a lowered to the normal specific gravity.

The blood-pressure *in the capillaries* is approximately measured by determining the amount of compression which is required to blanch a portion of flushed skin, *i.e.* to empty its capillaries; 2 centimeters of mercury is an average value.

The blood-pressure beyond the capillaries, *i.e. in the veins*, is exceedingly small and variable; in the jugular vein it is about  $\frac{1}{2}$  centimeter above and below zero in expiration and in inspiration respectively. There is danger of the entrance of air into a vein in operations at the root of the neck, owing to this negative pressure during inspiration.

The blood-pressure *in the heart* itself differs in the different chambers and oscillates with systole and diastole; these oscillations succeed each other so rapidly that the mean pressure as recorded by a mercurial manometer is considerably lower than the real maximum within the ventricle during systole. By means of a manometer guarded by a valve arranged so as to prevent the column of mercury from falling after having been raised by systolic pressure, Goltz and Gaule determined the maximum intraventricular pressure, *i.e.* that obtaining during ventricular systole; on reversing the valve, so that the column could fall but could not rise, they determined the minimum intraventricular pressure, *i.e.* that obtaining during ventricular diastole, and found that this minimum pressure was negative—in other words, that the ventricles have a suction action in the diastole. The following pressures were observed:—

	Maximum in systole	Minimum in diastole
Left ventricle . . . .	+ 15 cm. Hg	– 5 cm. Hg
Right ventricle . . . .	+ 6 cm. „	– 1·5 cm. „

By means of improved manometers, in which inertia is reduced to a minimum by the substitution of a spring for the heavy mercury, records of intraventricular pressure have been obtained that may be considered as accurate representations of its variations in the several phases of the beat. Fig. 23 is

an example of such a record; it shows a positive systolic followed by a slight negative diastolic pressure. In the auricles, except during their systole, the pressure does not

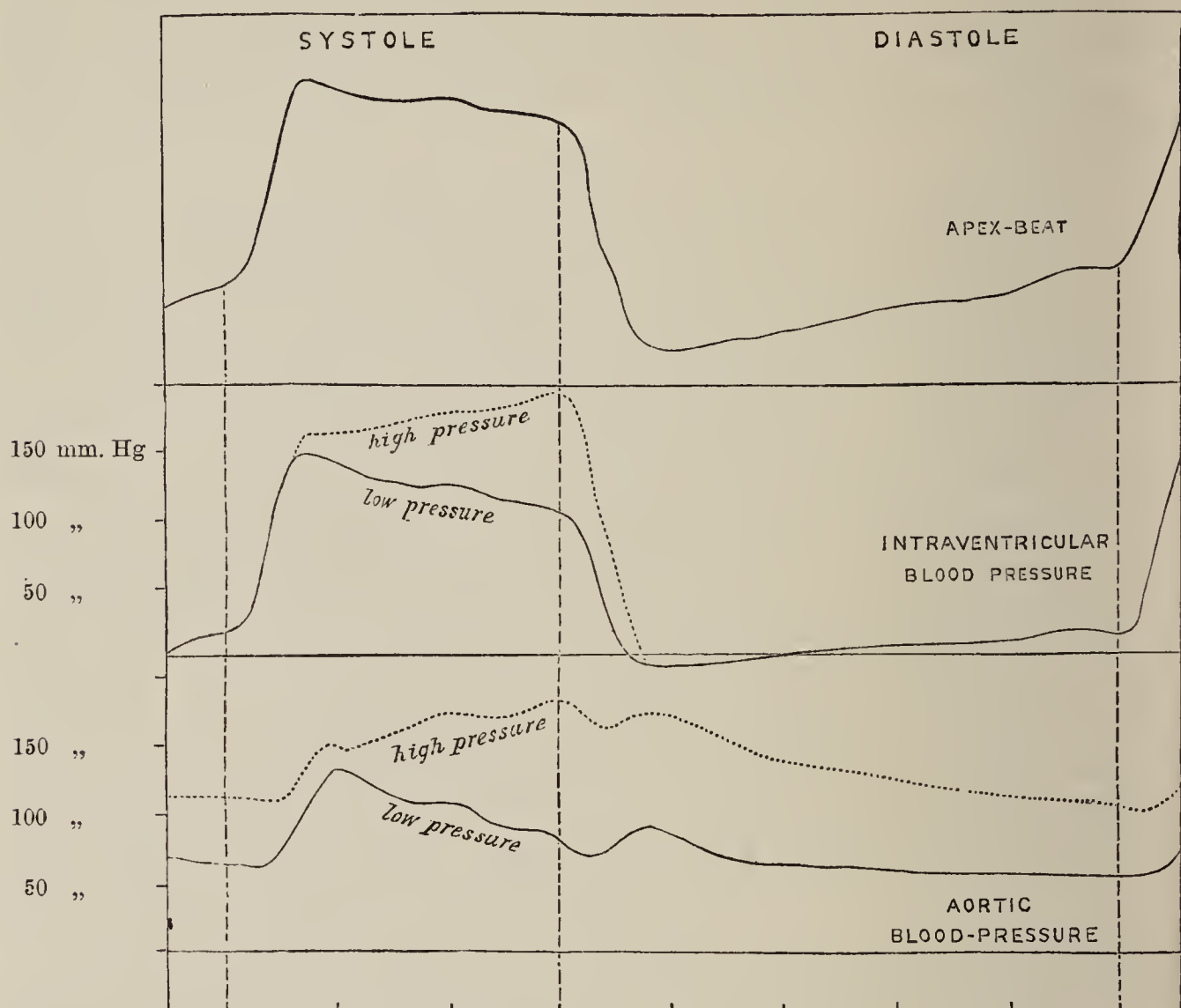


FIG. 23.—DIAGRAM AFTER HÜRTHLE (constructed from direct observations on the dog) TO ILLUSTRATE SYSTOLIC AND DIASTOLIC PRESSURE-VARIATIONS OF THE VENTRICLE AND AORTA ON MAN.

The noteworthy points of this tracing are: the similarity between apex-beat and intraventricular pressure; the post-systolic negative pressure in the ventricle; the higher pressure in the ventricle than in the aorta during systole; the unvarying duration of systole at high and at low blood-pressures; maintained pressure in the aorta; the negative pressure (diastolic notch) immediately followed by the diastolic wave. At 'low pressure' the systolic pressure in the aorta is double the diastolic pressure; in the high-pressure curve (dotted lines) the difference is proportionally smaller, *i.e.* a systolic augmentation of  $\frac{1}{4}$  to  $\frac{1}{3}$  of the mean arterial pressure. Notice the similarity between the intraventricular and aortic pressure-curves of the dog and the extravascular curves of apex-beat and carotid artery on man given in fig. 45, p. 87. Notice also that the systolic pressure augments from beginning to end at high pressure, and falls from beginning to end at low pressure.

exceed that in the large veins; hence even a small pressure within the pericardium will cause the auricles to collapse. The maximum systolic pressure in the left and right auricles amounts to about 5 and 2 centimeters of mercury respectively. Obstruction to the arterial circulation, as by compression of the aorta



or by vasomotor spasm, may raise the pressure in the left auricle to such an extent that it ceases to contract (w.).

*Influence of varying arterial pressure upon venous and upon pulmonary pressure.*—As might be expected, the general venous pressure has been found to rise with a fall of arterial pressure, and to fall with a rise. The mechanism of the relation is as follows: (1) if the fall of arterial pressure is due to reduced action of the heart (*e.g.* by vagus excitation), blood accumulates in the great veins, and venous pressure rises; (2) if the fall is due to dilatation of systemic arterioles, blood passes with greater facility into the veins, and venous pressure rises; (3) if arterial pressure is raised by increased action of the heart (*e.g.* after section of vagi), blood must be taken from the great veins with increased rapidity, and venous pressure will fall; (4) if a raised pressure is caused by constriction of arterioles, less blood passes into the veins, and venous pressure falls.

There are, however, exceptions to this rule; the effects on venous pressure of a dilated or contracted outlet of the arterial system may be masked by the effects of the increased or diminished capacity of the arteries caused by their diminished or increased tonus; if the medulla oblongata be destroyed or the splanchnic nerves be cut, venous as well as arterial pressure will fall, because blood accumulates in the relaxed abdominal vessels; if the medulla or splanchnic nerves be stimulated or the abdomen compressed, venous and arterial pressures will be raised, because blood is squeezed from the abdominal vessels into the rest of the vascular system. It is also obvious that a fall or rise of pressure, caused by hæmorrhage or by injection, will, if effectual at all, lower or raise venous with arterial pressure. Apart from these exceptions, the general rule holds good, that venous pressure falls with a rise and rises with a fall of arterial pressure. We shall find this relation illustrated in the alterations of pressure that accompany the movements of respiration; it is also well shown by 'Traube-Hering' curves, the pressure falling and rising in the veins as it rises and falls in the arteries (pp. 144, 148).

Alterations of pressure in the systemic vessels will influence the pressure in the pulmonary vessels in the same sense as in the systemic veins, and in a similar manner, as a consequence of greater or smaller onward flow; pressure in the pulmonary circuit will thus rise with a fall, and fall with a rise of aortic pressure. These effects are, however, very much reduced by the great



distensibility of the pulmonary vessels, which can accommodate a large addition of fluid without sensible rise of pressure, and which will permit the passage of an undiminished volume of blood, when the vascular channels have been experimentally reduced to less than one-half their sectional area (Lichtheim). It is to be observed, moreover, that a rise or a fall in pulmonary blood-pressure, due to free or limited supply of blood from the systemic veins, due in turn to free or limited arterial outlet, may be counterbalanced, or even reversed, by the retrograde effect of diminished or increased systemic obstruction; greater aortic pressure may in this way cause a rise in place of a fall of pulmonary blood-pressure. In sum, the blood-pressure and the resistance in the pulmonary circuit are low and constant, and very little influenced by variations of aortic pressure.

The converse relationship, *i.e.* the effect of varying pulmonary pressure upon aortic pressure, is of more importance. We shall find in studying the effects of respiration upon circulation, that alterations of pressure in the air passages, by pressing upon or by expanding the pulmonary vessels, can drive on the blood, or obstruct its passage, and so lead to a rise or to a fall of aortic blood-pressure. This is particularly the case in laboured respiration, and is illustrated in an extreme degree by Müller's and by Valsalva's experiments (p. 146).

**Further physical considerations.**—A drop of water starting from the level A will, by the time it has fallen through the space H to the level B, have a velocity  $= \sqrt{2 G H}$ . (G represents the acceleration per second under the influence of gravity, and is equal to 32 feet or 9.8 meters.)

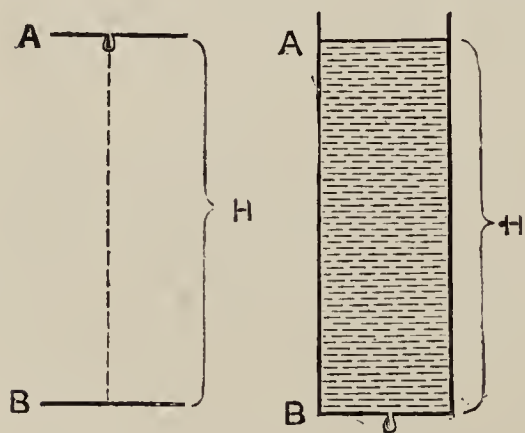


FIG. 24.

A drop of water starting from an orifice in B of a vessel of water filled up to A will have at B the same velocity as if it had fallen freely from A, viz.  $\sqrt{2 G H}$ . This is Torricelli's theorem.

Water escaping along a horizontal tube of uniform calibre from a reservoir in which a constant level is maintained would travel with the uniform velocity of  $\sqrt{2 G H}$  at all parts of the tube, *if there were no resistance*. But there is resistance at the inlet, along the tube, and at the outlet, in consequence of which the flow is retarded and a lateral pressure is maintained within the tube. This lateral or resistance pressure is indicated by the height to which water will rise in the vertical tubes 1, 2, 3 (fig. 25). Omitting, for the sake of simplicity, the

resistances at inlet and outlet, it will be seen that the levels in the three vertical tubes will diminish uniformly; the level in 1 indicating the lateral pressure produced by resistance in the horizontal tube from 1 to outlet; the level in 2 that produced by resistance from 2 to outlet, &c.

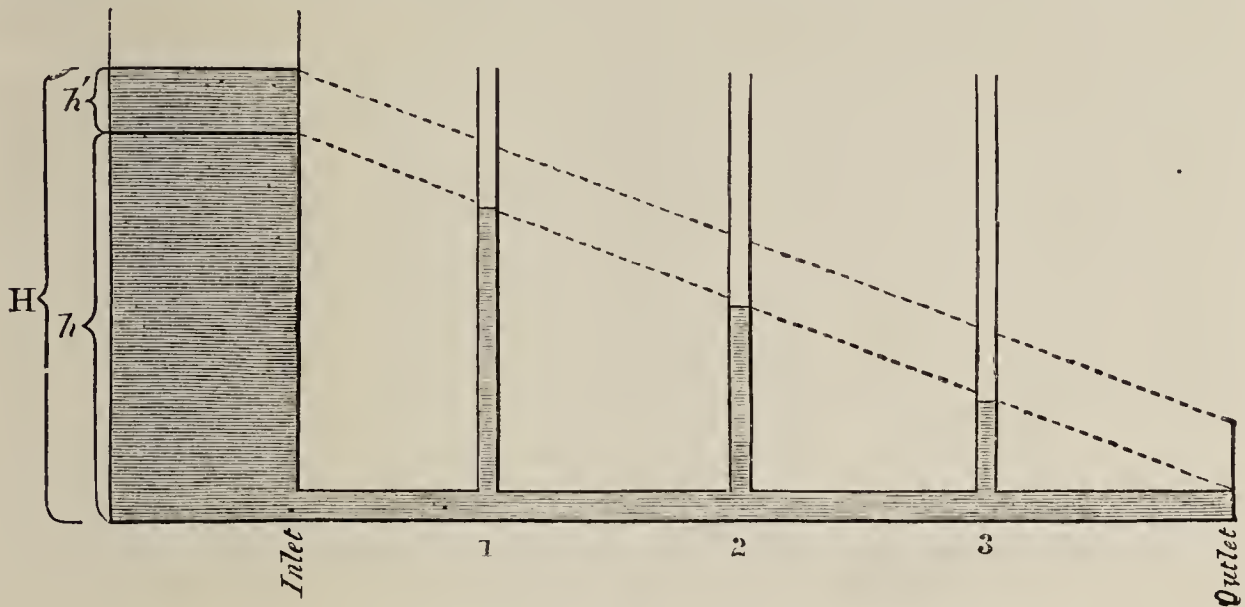


FIG. 25.

The total height or pressure,  $H$ , is divisible into two parts,  $h$  and  $h'$ ,  $h$  being the pressure due to resistance,  $h'$  being the pressure producing flow, or the 'velocity-pressure.' The mean velocity, which is uniform, is ascertained by experiment; its value per second is equal to

$$\frac{\text{Volume of outflow}}{\text{Time in seconds} \times \text{sectional area}} \quad \text{or} \quad \frac{Q}{t \pi r^2}$$

From the ascertained velocity the velocity-pressure  $h'$  is calculated by

$$V = \sqrt{2Gh'} \quad \text{or} \quad h' = \frac{V^2}{2G}$$

Water ejected from a syringe (fig. 26) with an internal pressure measured by the height of the column,  $H$ , would have the velocity  $\sqrt{2GH}$  in the absence of any resistance. But the velocity as observed would be much less, owing to resistance at the inlet  $i$ , along the tube  $t$ , and at the outlet  $o$ . The total pressure,  $H$ , is divisible into two parts,  $h$  and  $h'$ ,  $h$  being expended upon resistance,  $h'$  upon onward movement or velocity. The actual value of the velocity is only  $\sqrt{2Gh'}$ .

Along a tube of varying calibre the mean velocity will be greater in narrow than in wide portions, viz. inversely proportional to the sectional area  $\pi r^2$ .

At any given section of a tube, the velocity will be greater in the axis than at the circumference of the current. The mean velocity is an average value less than the greatest velocity in the axis, greater than the smallest velocity at the circumference.

Transferring these considerations to the heart and blood-vessels,



we recognise that the total cardiac pressure expends itself, 1st, in overcoming resistance ( $h$ ), 2ndly, in driving on the blood ( $h'$ ). The value of  $h$  is given by a manometer placed in connection with a large artery; <sup>1</sup> the value of  $h'$  is calculated from the observed velocity by the formula

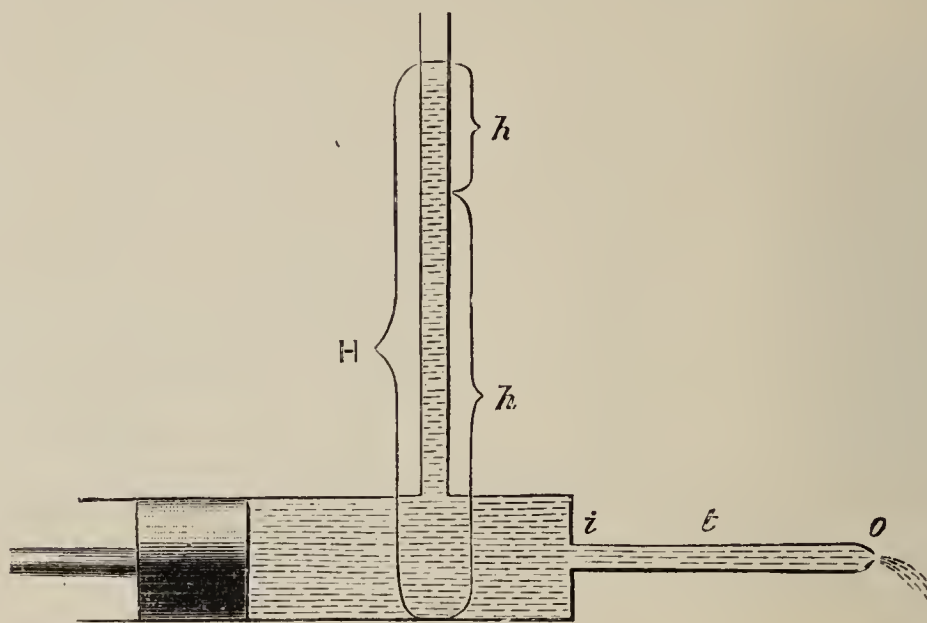


FIG. 26.

$h' = \frac{V^2}{2G}$ ; e.g. the velocity in a carotid artery having been observed to be 30 cm. per second, the value of the velocity-pressure,  $h'$ , will be  $\frac{30 \times 30}{1960}$  or .46 cm. H<sub>2</sub>O or .035 cm. Hg. Taking the lateral blood-pressure at 14 cm. Hg,  $h'$  is seen to be only  $\frac{1}{400}$  of  $h$ . Thus of the total cardiac pressure,  $H$ , on large mammals, and therefore presumably on man, by far the larger part constitutes the portion  $h$  by which resistance is overcome, the portion  $h'$ , by which blood is driven onwards, being only  $\frac{1}{400}$  part of the lateral pressure,  $h$ , or of the total pressure,  $H$ . The actual velocity in a large artery under average conditions of pressure and resistance is only  $\frac{1}{20}$  of the value that it would have with the same pressure, but with zero resistance; the theoretical value of the velocity in the absence of resistance is  $\sqrt{2GH}$ , the actual value in a large vessel is about  $\frac{1}{20} \sqrt{2GH}$ . The recognition of this fact is of practical importance: the velocity in an unopened carotid is about 30 cm. per second, whereas in a freely divided artery, with the resistance reduced almost to zero, it will (omitting any correction for reduced pressure) approximate towards the value 600 cm. per second.

<sup>1</sup> It appears at first sight as if the total pressure,  $H$ , is measured when a simple cannula is placed in the cardiac end of a divided artery, and the resistance-pressure,  $h$  only, when a T-shaped cannula is placed in the course of an artery. In the first case there is no current in the vessel; in the second, blood is flowing as usual; but practically there is no distinction between the two methods, which both give the resistance-pressure,  $h$ . With the simple cannula in the cardiac end of an artery, the whole artery is equivalent to a lateral tube springing from a vessel in which blood is flowing. Besides, as stated in the text, the difference between a total pressure,  $H$ , and a lateral pressure,  $h$ , in a normal artery, is so small as to be negligible.

It is not possible to obtain the normal current velocity in an artery by measurement of the quantity of blood escaping from its cut end in a given time. From such measurement it would indeed be possible to estimate velocity in the divided artery, but this would give no information as to the normal velocity in the undivided vessel. The velocity of current in a divided carotid, which we have seen is theoretically twenty times as great as in the undivided vessel, is actually five to ten times as great, owing to the fact that resistance is not reduced to zero, nor pressure fully maintained. From this consideration we may realise with what rapidity the blood will escape when a large artery is divided. Taking the sectional area of the carotid artery as  $\cdot 8$  square centimeter, the amount of blood passing through the normal vessel will be  $30 \times \cdot 8$ , or 24 c.c. per second, but hæmorrhage from the divided vessel would be at the rate of 120 to 240 c.c. per second. The entire amount of blood in a man's body is about 5,000 c.c.; the loss of half that amount is fatal; the time of grace after accidental division of a large artery, during which life can be saved, is therefore to be counted in seconds, and even from a small artery blood escapes with alarming rapidity.

In the veins the conditions are different; the resistance is small, the lateral or resistance pressure is low; the velocity of outflow from a divided vein scarcely, if at all, exceeds the velocity of flow within an undivided vein. On the other hand, comparing the velocities in the undivided vein, and in the undivided artery, no great difference is found—*e.g.* in the carotid the velocity is about 30 cm., in the jugular it is about 20 cm. per sec.

The relation between velocity and pressure differs in large tubes and in small tubes (*i.e.* less than 1 mm. diameter). In large tubes the velocity of stream varies as the square root of the pressure; *i.e.* with pressures of 1, 4, 9, 16, the corresponding velocity will be 1, 2, 3, 4. In small tubes the velocity of stream varies as the pressure; *i.e.* with pressures of 1, 2, 3, 4, the corresponding velocities will be 1, 2, 3, 4; moreover in small tubes variations of diameter sensibly influence the velocity, while in large tubes such variations have no effect; with constant pressure the velocity in small tubes varies directly as the radius squared; *e.g.* if blood runs at the rate of 1 mm. per sec. through a capillary with a radius of  $5 \mu$ , it will run at the rates 4, 9, 16, 25, 36 mm. per sec. in vessels with radii of 10, 15, 20, 25,  $30 \mu$ ; thus the pressure in a network of minute vessels being uniform, the blood will run much faster through the largest than through the smallest channels.

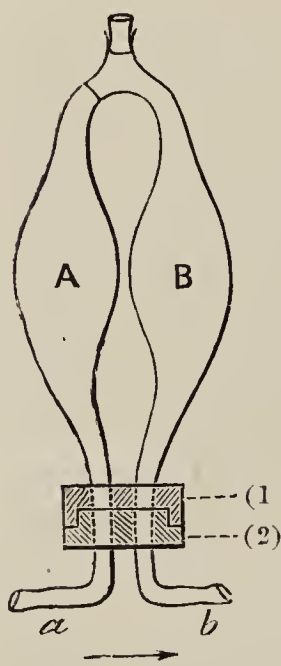
The average velocity of the blood-flow through the capillaries is  $\frac{1}{2}$  to 1 mm. per second; this very low velocity, as compared with the velocities in the carotid and jugular vessels, is owing to the much larger sectional area of the total capillary bed as compared with the sectional area of arteries or veins near to the heart, and to the much



greater lateral friction which the blood encounters in the minute channels.

In the living body the relation between pressure and velocity is variable and complex, and chiefly dependent upon variations of peripheral resistance; as we have already seen, greater resistance produces greater pressure with smaller velocity, smaller resistance produces smaller pressure with greater velocity. Moreover, the outflow from the large arteries, and therefore the velocity within them, depend upon the minute arteries and capillaries, which govern pressure and flow in the larger pipes; *cæteris paribus*, therefore, the velocity of arterial blood-flow must vary as pressure, and not as square root of pressure; thus an organ without variation in the diameter of its arterioles will be supplied by double or by half as much blood if the general blood-pressure rises to double or falls to half its normal value; if, with the rise of pressure, the arterioles of the organ itself should dilate, the organ will be still more copiously supplied; or if, with fall of general pressure, the arterioles contract, supply will be still further reduced. Velocity of arterial or venous blood-flow may be ascertained by direct observation; quantity of blood flowing through an artery may be calculated from the measured velocity and sectional area.<sup>1</sup> Thus, given the velocity of current in a carotid artery as 30 cm., the diameter of the vessel as 1 cm., we shall have the quantity of blood flowing through it equal to  $30 \times 3.14 \times \frac{1}{4}$ , or 23.55 c.c. per second. Or again, taking the collective sectional area of the feeding arteries of the brain at 30 square milli-

FIG. 27.—LUDWIG'S STROMUHR.



Two bulbs, A and B, fixed to the upper metal disc (1); two cannulae, *a*, *b*, fixed to the lower metal disc (2); the upper disc movable round the lower, so that the connection between the bulbs A B and the cannulae *a b* can be reversed, or interrupted. Cannula *a* fixed in central end of carotid, cannula *b* in peripheral end: bulb A filled with oil, bulb B with defibrinated blood. Blood from the central end of artery enters A, drives oil over to B, from which the defibrinated blood is driven into the peripheral end of artery. A being full of fresh blood, B is full of oil: the position of the bulbs is suddenly reversed by a half-revolution of the upper disc. B (full of oil) is now connected with *a*; A (full of blood) is connected with *b*. The manœuvre is repeated several times.

Given the capacity of a bulb, the number of times it has been filled and emptied, and the sectional area of the artery, the *velocity* of the blood-current is calculated—*e.g.* a stromuhr placed on a carotid artery of a small dog showed a flow of 90 c.c. per minute, *i.e.* 1.5 c.c. per second. The sectional area of the vessel was 5 square mm., *i.e.*  $\frac{1}{20}$  square cm. The rapidity of the current is, then, 1.5 divided by  $\frac{1}{20}$ , *i.e.* 30 cm. per second.

meters, we should find, as the amount of blood received by the brain per second, a quantity of about 9 c.c.

<sup>1</sup> To calculate the sectional area of a vessel, measure its diameter, square half this number, and multiply by 3.14—*i.e.* sectional area = radius<sup>2</sup> ×  $\pi$ .

The sectional area of a distended vessel is greater than that of an empty vessel

Or proceeding otherwise, the quantity of blood passing through an artery may be ascertained by direct observation ; velocity is then calculated from the measured quantity and sectional area of the artery employed.

Such measurements have been made by Volkmann, Vierordt, Chauveau, Ludwig, and by many others, and the chief instruments used have been the *hæmotachometer* of Vierordt, the *hæmodromometers* of Volkmann and of Chauveau, the *stromuhr* of Ludwig. The principle upon which the determination depends in the instruments of Vierordt

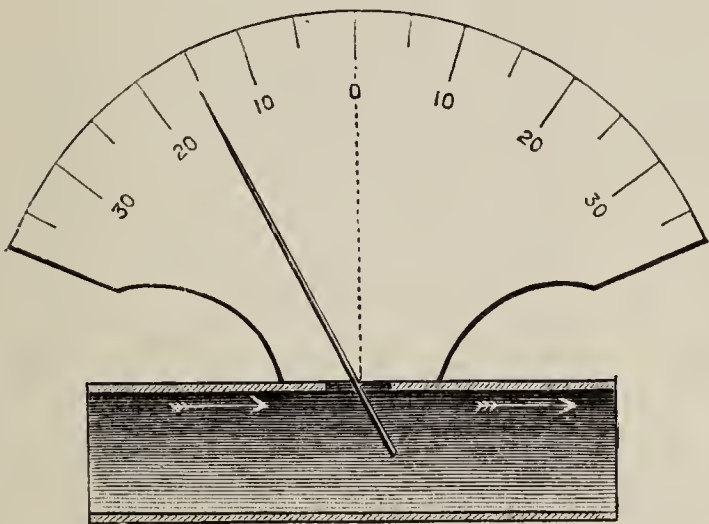


FIG. 28.—CHAUVEAU'S HÆMODROMOMETER.

and of Chauveau is the greater or smaller deflection of a pendulum by the more or less rapid stream of blood. By Ludwig's *stromuhr* the actual amount of blood passing through an artery in a given time is measured. The rate of current in the vessel is calculated from this amount, taking into account the sectional area of the artery employed.

The *stromuhr* is best adapted for giving information concerning the total amount of blood traversing an artery. The dromometer —*e.g.* the carotid artery at a distension-pressure of 20 cm. of Hg has about twice the sectional area of the same vessel at zero-pressure.

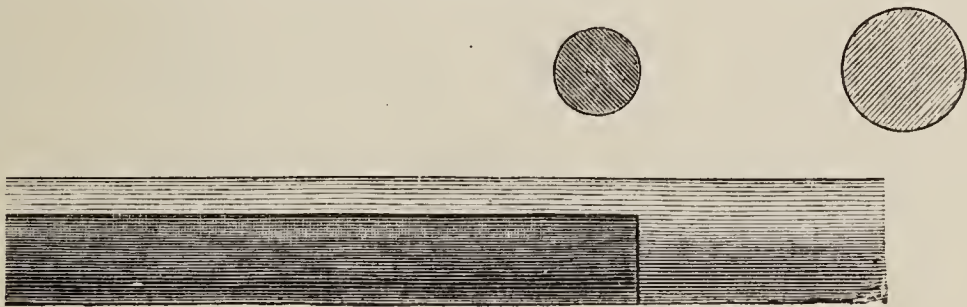


FIG. 29.

Portion of a human carotid artery at zero-pressure (dark shading), and distended by a pressure of 20 cm. Hg (light shading).

	At 0	At 20 cm.
Capacity . . . . .	1·9 c.c.	5·46 c.c.
Length . . . . .	50 mm.	70 mm.
Diameter . . . . .	7 „	10 „
Sectional area . . . . .	38 sq. mm.	78 sq. mm.



or dromograph shows the rapid variations of blood-flow which accompany rapid variations of pressure. A better instrument in use for this purpose is based upon the difference of pressure produced in 'Pitot's tubes.' In fig. 30, 1, 2, and 3, are Pitot's tubes in a long pipe leading from the bottom of the reservoir R. If there is no resistance at the outlet the water will stand at the levels  $a, a, a$  in the three tubes; if the outlet is closed the water will stand at the levels  $c, c, c$ ; if the outlet is partially open the water will stand at some such level as  $b, b, b$ . With a free outlet and maximum outflow, the difference

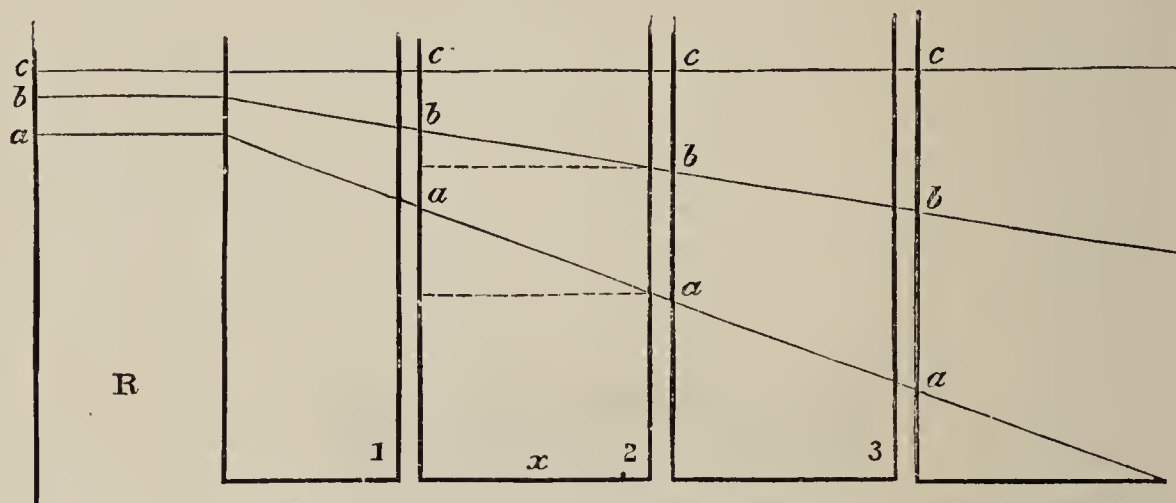


FIG. 30.

between any two levels  $a, a$ , is at a maximum; with a closed outlet and no outflow there is no difference between any two levels  $c, c$ . Between these two extremes it is obvious that the difference of level between any two tubes,  $b, b$ , will be greater or smaller as the outflow is greater or smaller, in other words, the difference indicates rate of flow. It is, moreover, obvious that any constriction, say at  $x$ , will increase the differences of water-level in the two tubes 1 and 2.

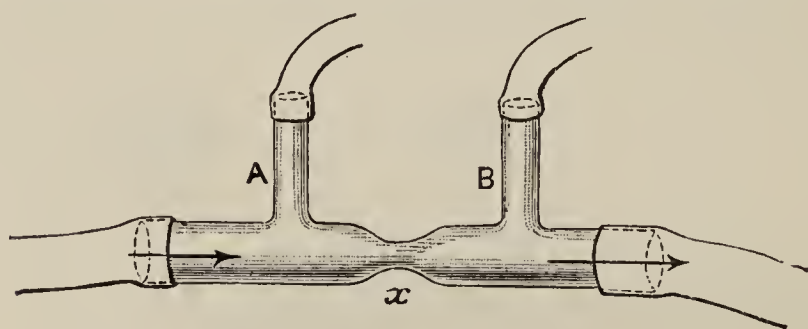


FIG. 31.

Variations of blood-flow can be determined upon this principle. A cannula (fig. 31) with two lateral branches, A, B, and an intermediate constriction,  $x$ , is fixed in the artery. Tubes from A and B lead to

two manometers. The level in the manometer connected with A will be higher than in that connected with B, and the difference will be greater or smaller according as the velocity is greater or smaller. If the two manometers be fixed in front of each other before a slit in a screen, the upper surfaces of the two columns can be photographed on a travelling sensitive surface, set up in a dark chamber behind the screen, and the varying difference between the two manometer levels thus recorded.

Alterations of velocity may be of central<sup>1</sup> or of peripheral origin; in either case they are indicated by the greater or smaller difference of pressure in A and B, but in the former case both pressures rise and fall with rise and fall of velocity, and in the latter both pressures rise and fall with fall and rise of velocity. Thus—

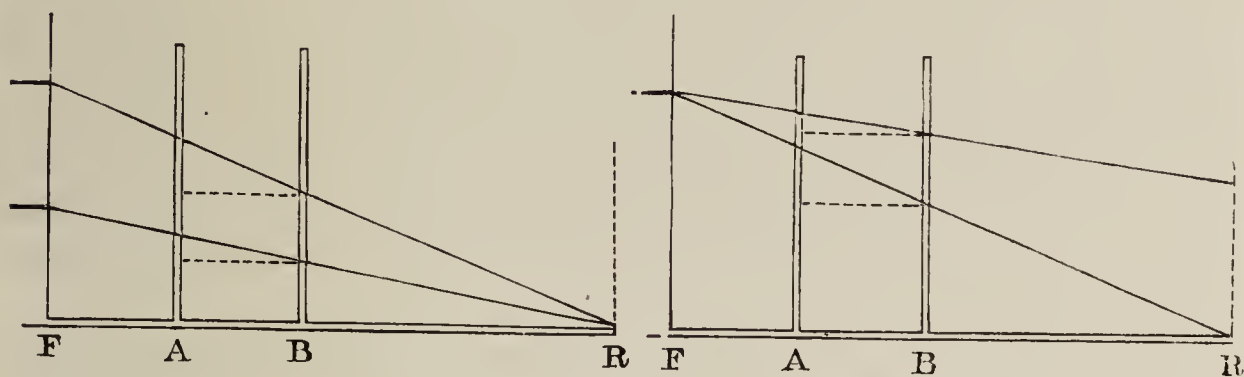


FIG. 32.

+ difference with + pressure indicates + velocity	} of central origin.
— difference with — pressure indicates — velocity	
+ difference with — pressure indicates + velocity	} of peripheral origin
— difference with + pressure indicates — velocity	

In the first two cases, the ratio between the pressures A and B remains unaltered; in the last two cases, the ratio of A to B increases with greater velocity (lower pressures), and diminishes with smaller velocity (higher pressures).

*The time of circulation*, *i.e.* the time occupied by one complete circuit of the whole mass of the blood, or—what practically amounts to the same thing—the time taken by any drop of blood to accomplish one complete circuit, is ascertained by injecting an easily recognisable salt (ferrocyanide of potassium, or preferably ferrocyanide of sodium) into the jugular vein of one side, and measuring the time that elapses before the salt is detected in the blood flowing out from the other jugular. During the interval between injection and detection of the salt, the latter has been conveyed with the blood through the right heart, pulmonary vessels, left heart and systemic vessels, *i.e.* through one entire circuit. Hering, by whom this method was first employed, found that the time is greater in large than in small animals, but that it corresponds very constantly with a definite number of heart-beats, *viz.* 28. Thus, for instance, in a horse with a pulse-frequency of 42 per minute, the time was 40 seconds; in a rabbit, with a pulse-frequency of 168 per minute, the time was only 10 seconds; *i.e.* in each

<sup>1</sup> 'Central,' that is to say, on the A side of the cannula, but not necessarily 'cardiac,' for evidently the pressure on the A side may be influenced by the peripheral state in districts other than that served through the cannula—*e.g.* vasoconstriction of the splanchnic area would cause rise of pressure and of velocity in the carotid artery. Practically, the relations described in the text could only be verified on large animals—*e.g.* on the carotid artery of the horse, greater velocity during mastication, smaller velocity during excitation of the sympathetic, due to dilatation or constriction in the peripheral distribution of the carotid.



case the circuit was accomplished with 28 heart-beats. Transferring this datum to the human subject with a pulse-frequency of 70 per minute we obtain 24 seconds as the time of circulation. The estimate so formed is obviously only an approximate one; several accessory factors influencing the actual results are necessarily neglected; thus the possible diffusion of the salt is not taken into consideration, nor the fact that different portions of the blood must traverse the circuit comparatively quickly by way of the head, and comparatively slowly by way of the lower extremities, kidneys and liver. Stewart finds, in rabbits, *e.g.*, that blood passes through the lungs in 2.5 seconds, round the head in 2 seconds, through the hind limb in 4 seconds, through the kidney in 8 seconds.

*The capacity of the heart's chambers.*—Measurements of the capacity of the auricles and of the ventricles taken after death are valueless or misleading, because the amount of fluid which these cavities will contain varies with their more or less contracted state, and is no measure of their working or functional capacity during life. We can, however, indirectly form such an estimate from various other data and considerations.

(1) Admitting as data, a total mass of blood in the body = 5,000 c.c., and 28 heart-beats required to send this mass once round the circuit, we obtain 178 c.c. as the amount which must be expelled by each ventricle at each contraction. This amount corresponds with the estimate generally quoted on the authority of Volkmann to the effect that each ventricle discharges at each beat an amount of blood equal to  $\frac{1}{400}$  the body-weight, *i.e.* in a man of 70 kilos each ventricle discharges 175 grammes of blood. But the estimate is probably too high. The number 28 expresses the quickest time in which ferrocyanide passes round the vascular system, and it is probable that the salt has made its appearance before the entire mass of blood has passed round. Howell and Donaldson estimate the ventricular discharge at  $\frac{1}{700}$  the body-weight, a fraction which is probably nearer the truth than  $\frac{1}{400}$ , *i.e.* in a man of 70 kilos each ventricle would expel at each beat 100 grammes, or nearly 4 ounces. The estimates of Zuntz and of Tigerstedt are lower still, viz. 50 to 100 c.c.

(2) A supplementary estimate, although not in itself of much weight, may be formed from other data. Assuming (1) the rate of the blood-current in the aorta = 30 cm. per second, (2) the sectional area of the aorta = 6 sq. cm., we must have flowing through the aorta 180 c.c. per second, which therefore would be the systolic discharge of the left ventricle with the heart beating at 60 per minute.

(3) An entirely different series of data can be utilised to form an estimate of the ventricular capacity. We shall find in the study of respiration that the average amount of oxygen absorbed is 375 c.c. per minute, and we may assume that arterialised blood returning from the lungs contains 5 c.c. per 100 more oxygen than venous blood. Under

these conditions the amount of blood by which the absorbed oxygen is carried off will be 7,500 c.c.; *i.e.* with the heart at 72 per minute, the right ventricle will discharge 104 c.c. of blood at each beat. We shall probably be not far from a true estimate by taking the average value of the discharge from each ventricle at each systole to be about 3 ounces, or 75 c.c.

It is obvious that the working capacities of the two ventricles must be equal, for if at each contraction one ventricle discharged more blood than the other, the systemic circuit would become emptied and the pulmonary circuit filled, or *vice versa*. It is also obvious that the working capacity of an auricle must be somewhat less than that of a ventricle, for the ventricle when filled contains overflow blood from the auricle as well as the blood that is discharged into it by the auricular contraction.

*Work of the heart.*—The ‘work done’ by a contracting muscle is expressed by the *height* to which a weight is raised. In the case of the heart, the weight raised is the amount of blood contained in the ventricles, the height to which that weight would be raised is the height of intraventricular blood-pressure during systole. From these data it is easy to calculate the ‘work’ done at each systole, and knowing the pulse-frequency, the average per hour or per day is also known. Admitting for the left ventricle an average discharge of 75 grammes, and a systolic pressure of 2 metres (about 15 centimetres Hg), the work done at each contraction will amount to 150 grammeters; to this amount we may add 50 grammeters as the work done by the right ventricle and by the two auricles, making up a total of 200 grammeters or  $\frac{1}{5}$  kilogrammeter as the work done by the heart at each beat. With a pulse-frequency of 72 per minute this would amount to 864 KgM. per hour, or more than 20,000 KgM. per diem. The whole of this energy is expended in the body, partly in overcoming resistance in the vascular system, and partly transformed into and discharged as heat; in this form the contractions of the heart yield about  $\frac{1}{50}$ th of the total daily heat-production.

*Distensibility of different vessels.*—The most distensible vessels are the pulmonary artery and its branches; the least distensible vessels are the systemic veins; the systemic arteries occupy an intermediate position in this respect, being less distensible than the pulmonary artery, more distensible than a systemic vein. The distensibility of an artery differs moreover from that of a systemic vein in this respect: a series of pressures of 1, 2, 3, 4, . . . &c., cm. Hg will cause on the artery a series of expansions at first increasing and then diminishing, on the vein a series of expansions which decrease from the first. According to Roy (who has shown that weighted strips of arteries behave in a similar manner) the turning-point, indicating the maximum of distensibility, coincides with the normal arterial pressure.

*Volume-recorders.*—A method which has been much used of late



years to study the circulation is that which is based upon measurement of the varying volume of parts or organs by means of the *plethysmograph* (Mosso) and the *oncometer* (Roy).<sup>1</sup> The plethysmograph is applicable to an arm or leg, or to a single finger; the part is enclosed in a rigid vessel full of water and communicating by a tube with an open vessel which is graduated or which contains a float. As the limb swells, water is driven into the graduated vessel, or raises the float; as the limb shrinks, water is drawn from the graduated vessel, or lets the float fall; when used as a recording instrument the float is connected with a light lever that traces its movements on a smoked cylinder. The oncometer is applicable to the spleen and kidney; the complete instrument is composed of two parts—the oncometer, which encloses the organ, and the oncograph, which records the movements of a piston; the two parts communicate by a tube, and the whole apparatus is filled with oil.

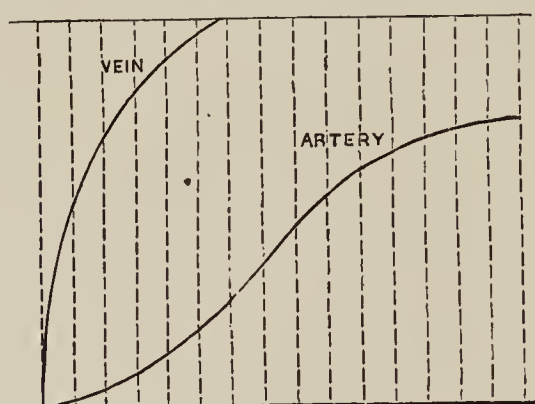


FIG. 33.

Capacity-curve of a vein distended by regularly increasing pressures from 1 to 60 mm.  $H_2O$ ; the same of an artery distended by pressures from 1 to 150 mm. Hg. (After Roy.)

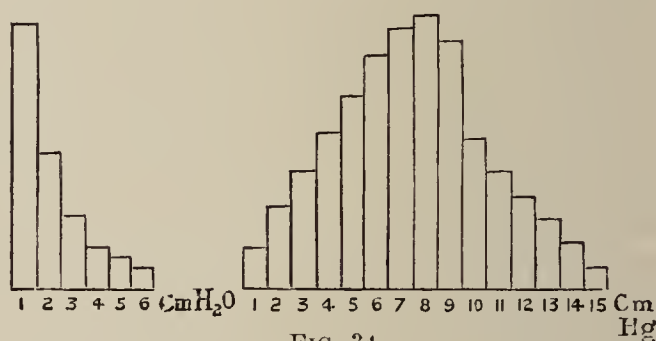


FIG. 34.

Increments of arterial and of venous capacity caused by equal increments of pressure. (After Roy.)

Variations of volume do not by themselves yield simple and unequivocal data, but require to be supplemented by readings of arterial and of venous blood-pressure. The variations in volume of a part or organ may be either active or passive; the volume of an arm, for instance, may be increased by dilatation of its arterioles or by obstruction of its veins; it may be diminished by constriction of its minute arteries, or by obstruction of its main artery. Generally speaking, the volume of a part or organ increases and diminishes with its functional activity, greater activity being accompanied by dilatation of the vessels, less activity or rest by constriction of the vessels. These are active variations effected by the vasomotor nerves of the part itself. But passive variations may also be produced, in consequence of variations of the general blood-pressure brought about by vasomotor nerves in other parts of the body. The kidney, for instance, may shrink or swell with a rise or with a fall of blood-pressure. If it shrinks with a rise or swells with a fall, these variations are local active changes due to

<sup>1</sup>  $\pi\lambda\eta\theta\acute{\upsilon}\omega$ , to swell;  $\acute{\upsilon}\gamma\kappa\omicron\varsigma$ , bulk.

constriction or to dilatation. If it shrinks with a fall and swells with a rise (as occurs after section of the renal nerves), these variations are local passive changes of an organ more or less distended by the general

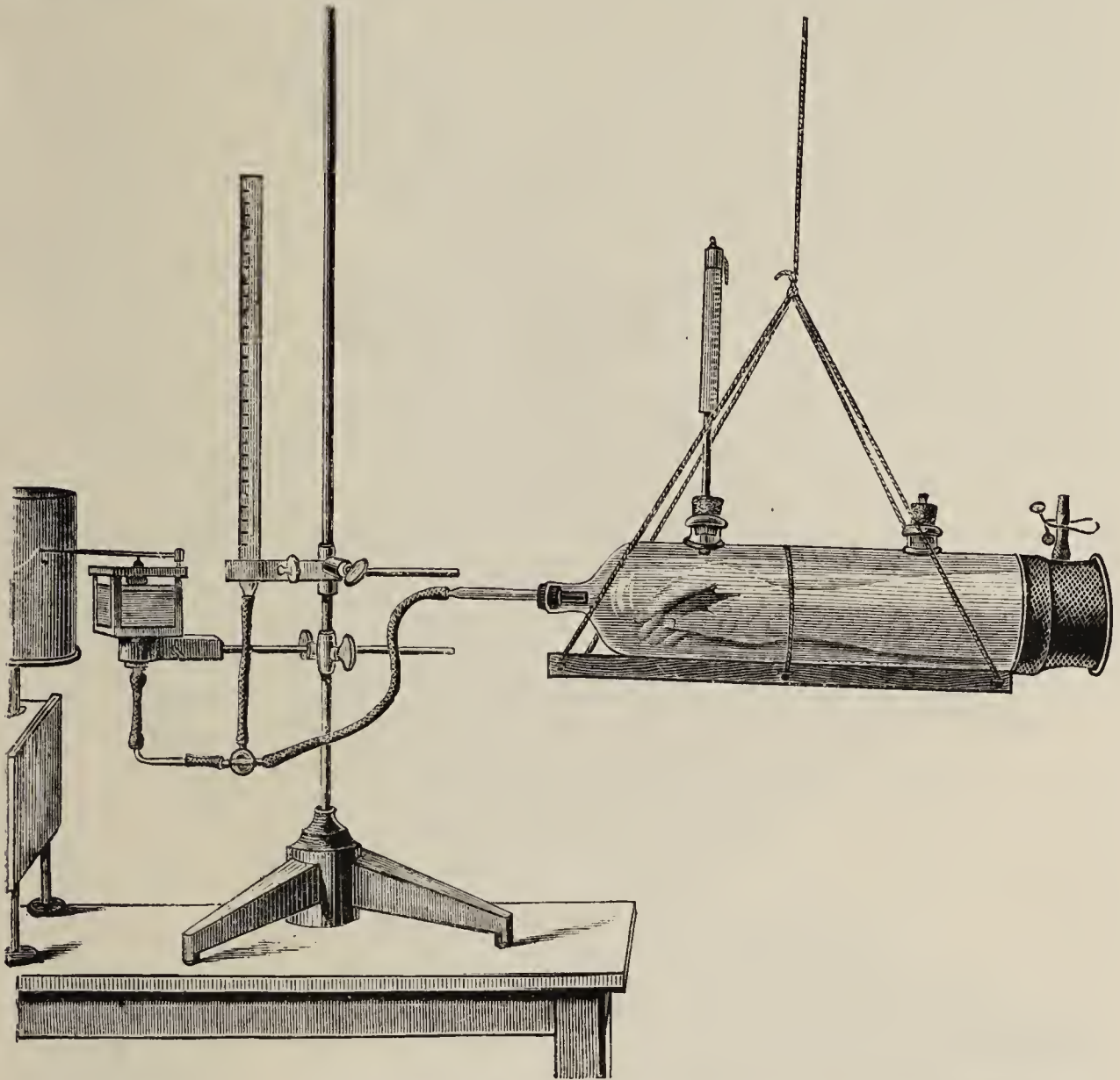


FIG. 35.-MOSSE'S PLETHYSMOGRAPH.

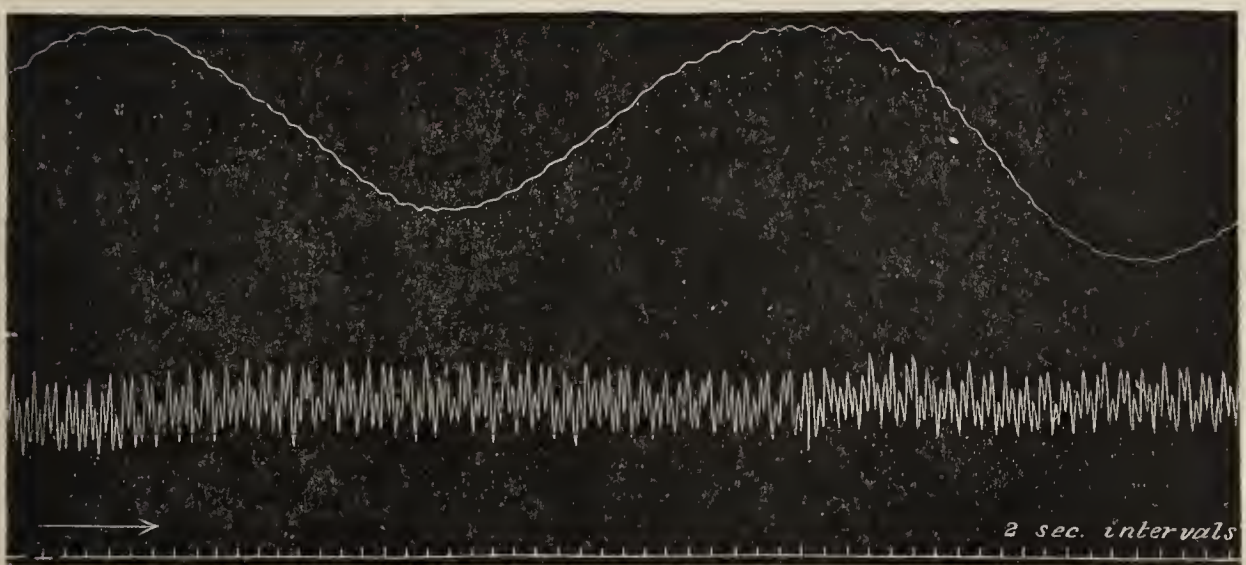


FIG. 36.

Simultaneous tracing of spleen volume (upper line) and of arterial pressure (lower line); rhythmical contractions of the spleen without sensible variations of arterial pressure. (Roy.)



blood-pressure. And when the nerves are intact the volume of the organ may either vary in the same sense as the general pressure, or in an opposite sense, according as the general pressure-changes overbear

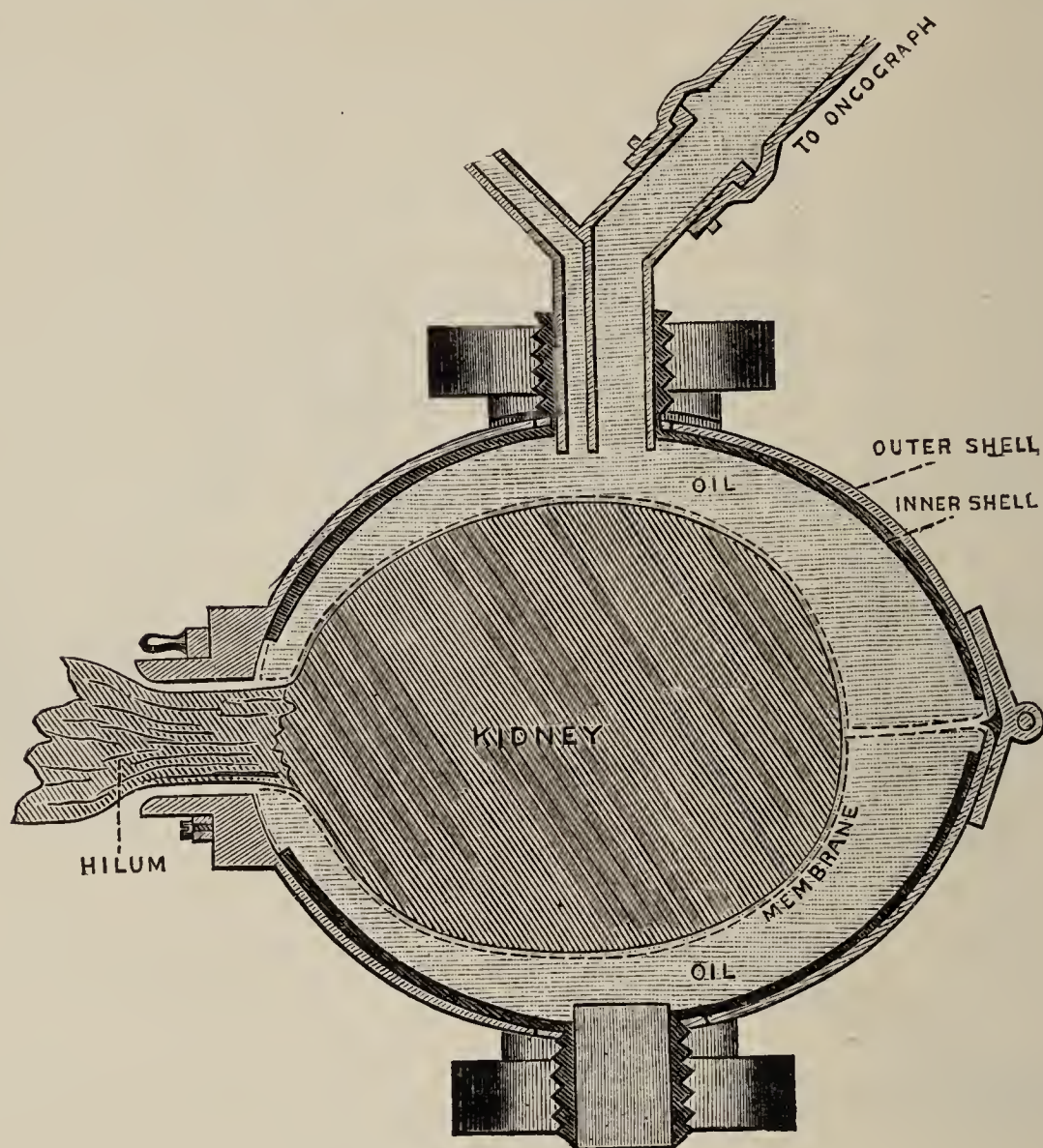


FIG. 37.—ROY'S ONCOMETER.

the local actions, or as the local actions exceed the general vasomotor changes of which they may form part. Other organs which are not directly influenced to any appreciable extent by vasomotor nerves, as, for instance, the brain, do not actively swell and shrink; the passive

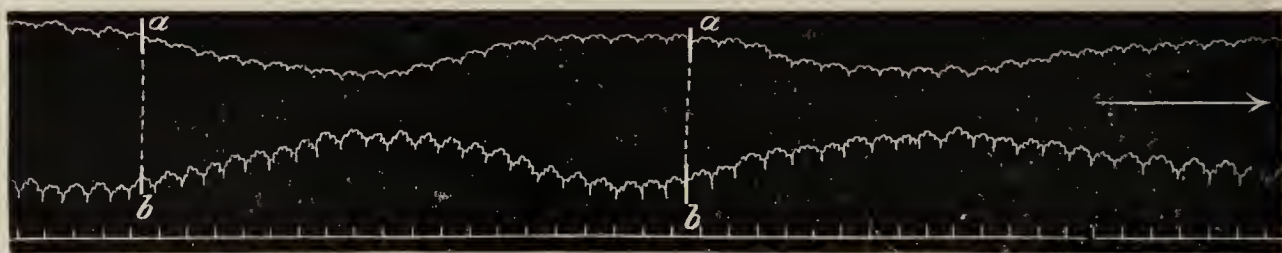


FIG. 38.

Simultaneous tracing of kidney volume (upper line) and of arterial pressure (lower line); the large undulations are Traube-Hering effects; with the rise and fall of arterial pressure the kidney volume falls and rises. (Roy.)

variations—distension by high blood-pressure, collapse of volume with low blood-pressure—are, on the contrary, well-marked and uncomplicated effects. The spleen, on the other hand, varies in most complicated



fashion ; it is a very distensible organ, and can therefore undergo considerable passive changes ; it is abundantly supplied with vasomotor nerves, and has therefore a considerable range of active vascular variations of volume ; and thirdly, it is highly contractile by virtue of its framework of involuntary muscle, so that, independently of passive and vasomotor variations, it can rhythmically dilate and contract. (Fig. 36.)

The increase of volume produced at each heart-beat is an indication of the accelerated blood-flow then occurring ; the venous outflow from the limb being constant, it is evident that a pulsatile increase of volume must be a measure of a pulsatile acceleration of arterial inflow (Fick, v. Kries).

The most notable of Mosso's plethysmograph experiments is that intended to demonstrate a variation of cerebral circulation coincident with cerebral exertion ; Mosso found that the volume of the arm was diminished during the performance of a calculation or other mental effort, and concluded that the effect was due to increased blood-supply to the brain ; the results of the experiment are, however, not sufficiently regular to bear out this conclusion ; alterations of volume of the arm, when they occur at all, are probably due to alterations of respiratory movements.

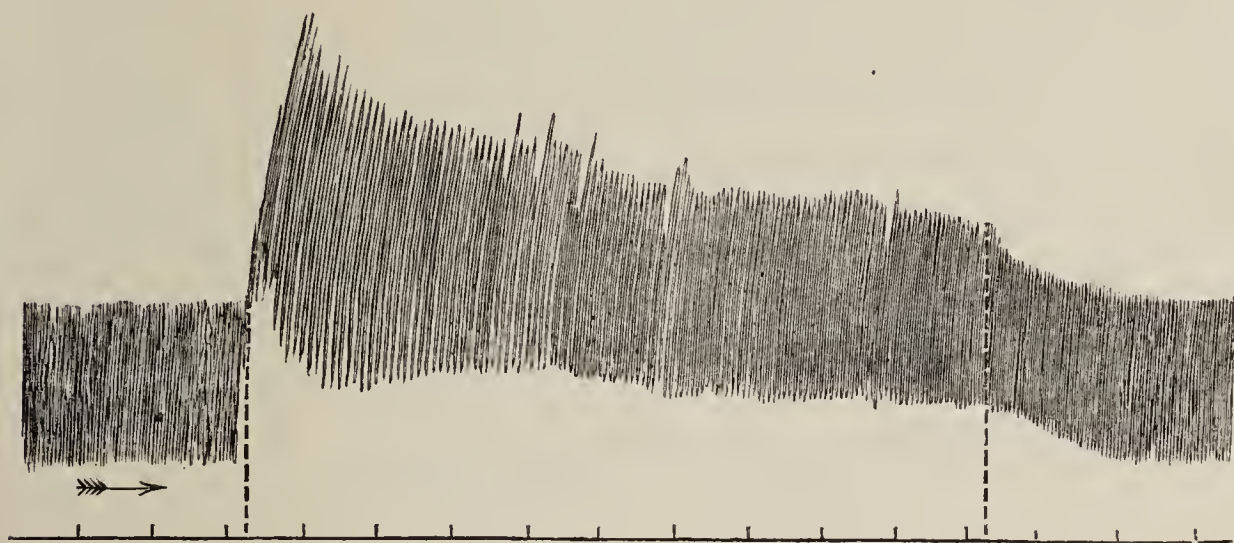


FIG. 39.—CARDIOMETER TRACING. (Roy and Adami.)

*The cardiometer.*—A volume-recorder on the same principle has also been adapted by Roy to the contracting heart, and by its means the output of blood per contraction or per minute may be measured, as well as the more or less distended state of the organ in various circumstances. Roy and Adami found, *e.g.*, that during vagus slowing the ventricles became distended, the 'output' per contraction increased, but the output per minute diminished. Another point of importance definitely ascertained by means of the cardiometer is that the ventricles do not become completely emptied at each contraction ; with high arterial pressure, as in asphyxia, the residual blood in the ventricles is at its greatest ; it should, however, be realised that the heart in a cardiometer is not under perfectly normal conditions.



The *sphygmomanometer* is an instrument by means of which the pulse-tension can be approximately measured on man : the principle upon which it is based being the determination of the amount of counter-pressure that is just sufficient to extinguish the pulse. A simple form of the instrument is as follows : an elastic finger-stall distended with air and connected with a mercury manometer is squeezed down upon the radial, or preferably the temporal, artery until the pulse is felt to vanish beyond the point of compression (recurrent pulsation in the radial being, if necessary, eliminated by compression of the ulnar artery) ; the pressure of air within the finger-stall as indicated by the manometer now just exceeds the blood-pressure within the artery during cardiac systole (w.).

The plethysmograph has also been employed for the same purpose, the principle of counter-pressure being utilised as follows : pressure of the fluid surrounding the limb (arm, hand, or finger) is raised until a point is reached at which the pulsatile alterations of volume are at a maximum ; this point is taken to indicate when the pressure of fluid on the limb is above the intra-arterial blood-pressure during cardiac diastole, but below it during cardiac systole.

**The pulse.**—At each beat of the heart about three ounces of blood (or 75 c.c.) are forced by the left ventricle into the aorta and added to the mass of blood which is being pressed onwards throughout the arterial system. This sudden addition to the contents of an arterial system which is already distended gives rise to a pressure-wave throughout all the arteries of the body, called the arterial pulse, which can be felt in any superficial artery. The most readily accessible artery for this purpose is the radial ; other accessible arteries are the temporal, carotid, axillary, brachial, femoral, popliteal, anterior and posterior tibials, in all of which the arterial pulse may be felt, and, if desired, examined by means of recording instruments. The practical value of the pulse is that it affords means of judging of the state of the circulation ; counting the pulse is the readiest means of ascertaining the frequency of the heart's beats, and the careful study of the tension of the pulse affords valuable information regarding the state of the arteries ; it enables us to judge whether the general arterial pressure is high or low or about normal. The normal *frequency* of the pulse (*i.e.* of the heart's beat) is about seventy per minute in the male, eighty per minute in the female, and still higher in children ; in the foetus before birth it is 130 to 140. The frequency is increased by exertion, by taking food, by alcoholic stimulants, by some forms of emotion, in fevers, and in all kinds of debilitating disease. It is diminished during rest,

during sleep, in some forms of emotion, in cerebral coma. As regards emotion, it is generally pleasurable and exciting emotion that raises the pulse-frequency, painful and depressing emotion that lowers it. As regards exertion, even the slight differences of exertion which insensibly accompany differences of posture affect the pulse-frequency; thus, for instance, in the same person the pulse has been counted in the lying posture 70, sitting 75, standing 80, and after a short run 120; in a weak or sick person the differences with posture may considerably exceed those observed in health; thus the difference in the sitting and lying postures may be twenty beats; it is therefore worth noting, when it is possible, what that difference is, for the number gives some measure of the weakness of the subject. Temperature also affects the pulse-frequency—high temperature raises it, low temperature lowers it; in the same person with a normal pulse-frequency of 72, the pulse has been counted in the hot chamber of the Turkish bath 96 per minute, and in a cold bath 60 per minute. Emotion from a purely imaginary and non-existent cause, is of itself sufficient to alter the pulse-frequency; in the hypnotic state in which vivid fictitious ideas can be easily provoked, such modifications are particularly well marked.

The pulse may feel *large* or it may feel *small*; these terms signify that the impression is conveyed to the observer of a large or of a small wave of blood passing under his fingers as they rest upon the artery; but the terms are apt to mislead, as will presently appear from other considerations. The same remark applies to the terms *strong* and *weak*, for an apparently 'strong' pulse is common when arterial pressure is low, an apparently 'weak' pulse when arterial pressure is high. It is more important to pay regard to the *compressibility* of the pulse and to speak of the *hard* pulse or the *soft* pulse. A hard pulse is one that requires considerable pressure of the fingers upon the artery to obliterate it; the artery feels distended between the beats, and the more the fingers compress it—within certain limits—the more forcible does the beat appear. A soft pulse is easily obliterated by compression, and appears most forcible when the fingers are lightly applied. The hard pulse is a sign of high arterial tension or pressure, the soft pulse is a sign of low arterial pressure; generally speaking, hard pulses are of lower frequency than soft pulses, and usually the hard pulse appears small and weak in comparison with the soft pulse if both are felt with a light touch. The terms *long* and *short* are sometimes used to



signify that each beat appears to last under the finger for a long or a short period ; they are unnecessary terms, for the long pulse is hard, the short pulse is soft. *Slow* and *quick* are sometimes used to mean the same as long and short respectively, and are equally unnecessary terms. *Rapid* and *slow* are occasionally employed to signify that the pulse is felt at a short or long interval after the heart's impulse. The supposed differences of the interval are imaginary ; there is an interval, it is true, but its variations are so slight as to be quite inappreciable without the aid of delicate instruments.

From these remarks it appears that the characters of the



FIG. 40.

Tracings of an irregular palpitating heart due to excessive tobacco-smoking (upper line), and of the same heart after administration of digitalis (lower line).

pulse that can be recognised and named without ambiguity are—(1) frequency, (2) compressibility. The pulse of high arterial tension is infrequent and hard (*rarus et durus*). The pulse of low arterial tension is frequent and soft, and sometimes palpably dicrotic (*frequens et mollis*). Every normal and most abnormal pulses are dicrotic in so far as their tracings reveal the presence of the dicrotic wave ; clinically, however, the term *dicrotic pulse* is reserved for cases in which the wave is so marked as to be easily felt ; a dicrotic pulse is a pulse of low tension, and when well marked gives to the finger of an observer a double tap with the rhythm of a postman's knock. An *intermittent* pulse occurs when, in the otherwise regular series of heart-beats, one or more beats are occasionally dropped, thus differing from an *irregular* pulse in which there are irregular intervals between the successive heart-beats, and usually great variety of strength of beat. The pulse of aortic regurgitation is so strikingly characteristic as to call for separate mention. It is termed a 'collap-

sing' or 'water-hammer' pulse, and its character is due to the fact that an abnormally large ventricle drives an abnormally large quantity of blood into an arterial system which, in consequence of backward escape through the aortic orifice, does not remain properly filled between the beats. In the examination of pulse-tension by the radial artery a *recurrent* pulse may often be felt beyond the point where the vessel is completely compressed; such a pulse is due to free anastomosis with the ulnar artery, and ceases when both radial and ulnar arteries are compressed (w.). A *venous* pulse may occur under various conditions; physiologically it is produced when the arterioles are widely dilated, so that the arterial pulse is propagated through them and through the capillaries into the veins; an instance of this kind of venous

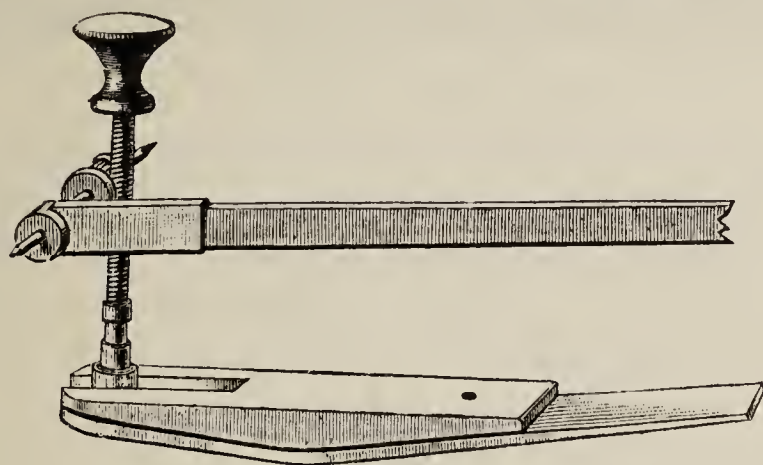
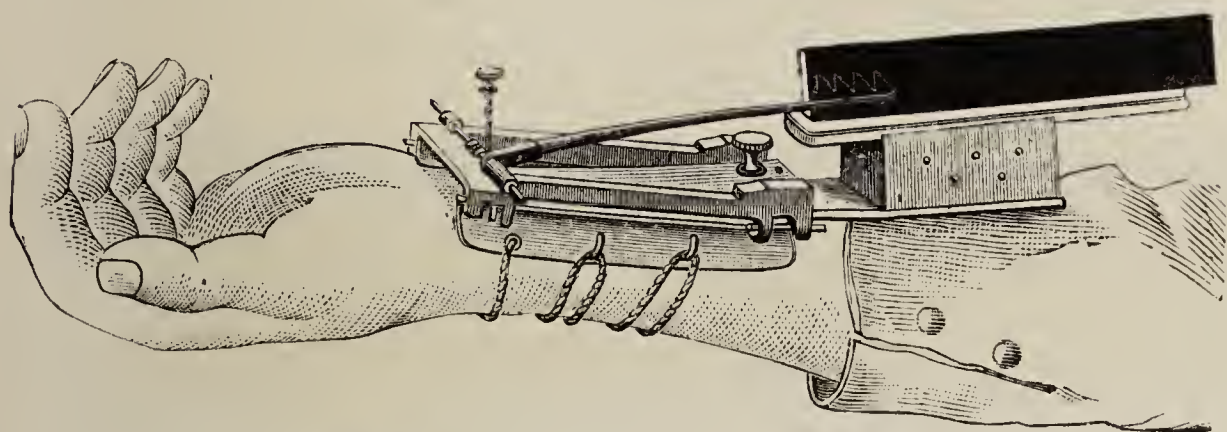


FIG. 41.

Marey's sphygmograph applied to the radial artery.  
Portion of lever and spring which presses upon the artery.

pulse occurs in the case of the submaxillary veins during excitation of the chorda tympani; it is also observable in the veins on the back of the hand after immersion in warm water. Another variety of venous pulse is the backward pulse observable in the jugular veins when from any cause the tricuspid orifice gapes during systole. A *capillary* pulse can be brought into evidence on most normal persons, and with great distinctness in cases of aortic regurgitation; it is seen as a systolic flush of colour at



the root of a finger-nail or in a portion of skin shortly after it has been compressed.

*The sphygmograph.*—If a lever rests upon an artery such as the radial at the wrist, so as to compress but not obliterate it, each pulsation will visibly raise the lever. A lever arranged thus and marking its movements upon a travelling surface constitutes a *sphygmograph*.

A normal pulse-tracing presents the following features: When the pulse-wave reaches the artery, where it is compressed by the lever, the compressed portion expands and the lever is raised, giving on the tracing the *primary* or *percussion* wave; the expansion of the artery having quickly reached its

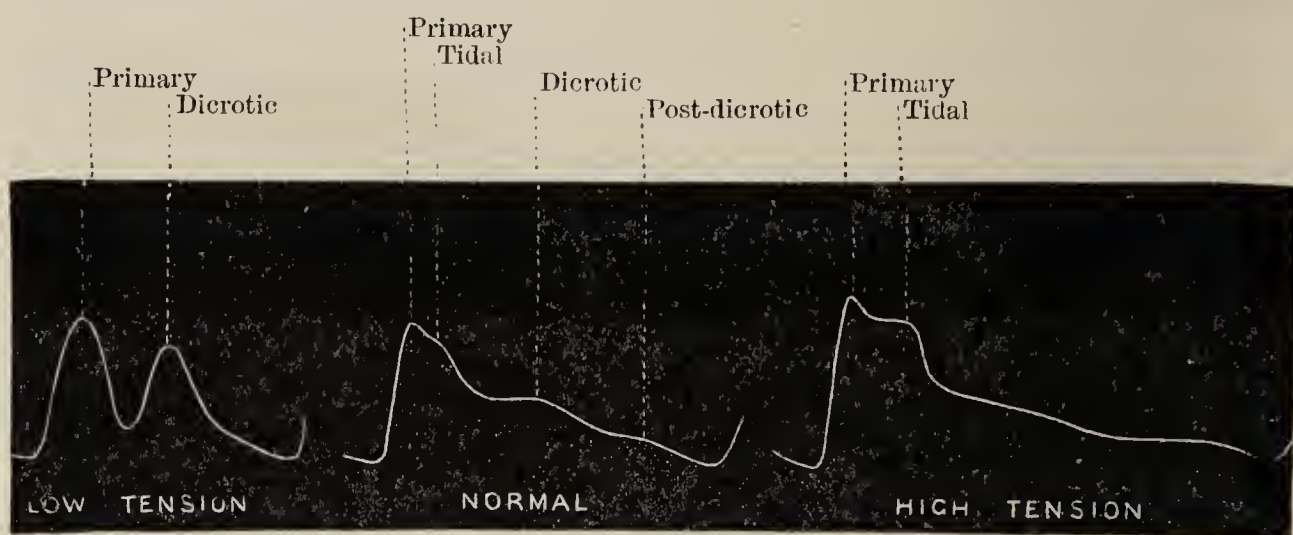


FIG. 42.

maximum begins to subside and the lever begins to fall; the decline is delayed by a brief period of maintained expansion, which may amount to an actual wave of renewed expansion; this interruption of the decline or second rise, as the case may be, is called the *tidal* or *pre-dicrotic* wave; the decline in the expansion of the artery now continues rapidly, and the lever falls correspondingly until the decline is suddenly arrested and replaced by a distinct increase of expansion constituting the *dicrotic* wave; between the two chief waves—the tidal and the dicrotic—a more or less distinct depression is marked, which is sometimes spoken of as the *dicrotic notch*. Subsequently to the dicrotic wave, the expansion of the artery continues to decline, the regularity of the decline being sometimes interrupted by a fourth wavelet, called from its position the *post-dicrotic* wave.

Are all these waves truly the expression of the varying states of pressure within the artery, or are they in part or wholly due to instrumental oscillations? To this question, which has been

much debated, we may answer at once that the percussion wave and the dicrotic wave are really caused by intra-arterial waves of pressure and consequent expansions of the compressed artery. If a horizontal jet from any small artery be received on a revolving drum covered with white paper, the blood marks a kind of pulse-tracing on the paper: such a tracing shows two waves, the primary wave and the dicrotic wave. These waves are therefore certainly arterial and not instrumental. The post-dicrotic wave is inconsiderable, it is probably instrumental, and it indicates nothing in particular. The pre-dicrotic or tidal wave is, properly speaking, due to a state of maintained expansion, rather than to any separate wave of expansion; it is probably exaggerated by the sphygmograph, the lever of which is apt to rebound after its fall from the primary or percussion rise. But apart from this possible source of deformation the tidal wave or state of maintained arterial expansion has a real value as an index of arterial condition. Normally in the radial artery it is not very pronounced; in the brachial or in the carotid artery—*i.e.* nearer the heart—it is much more marked; abnormally, it may be a well-marked feature of the radial pulse curve, and is then a sign that arterial tension is high, or that the arteries are abnormally rigid; it is a constant feature of the radial pulse-tracing taken on old people, and if it is found on the pulse-tracings of people at or below middle age, it indicates a prematurely ‘aged’ and rigid state of their arteries. Such a pulse would feel long and hard—*tardus et durus*; if due to high arterial tension it would be felt full between the beats, if due to rigid arteries it would be felt empty between the beats. When arterial tension is low, the tidal wave does not appear in the radial pulse-tracing, and may even be absent from the brachial tracing. The dicrotic wave is on the contrary exaggerated—so much so that the pulse can be felt to be double—whereas in high-tension pulses it is very ill-marked. The low-tension pulse is short and ‘compressible,’ and feels empty between the beats—*celer et mollis*.

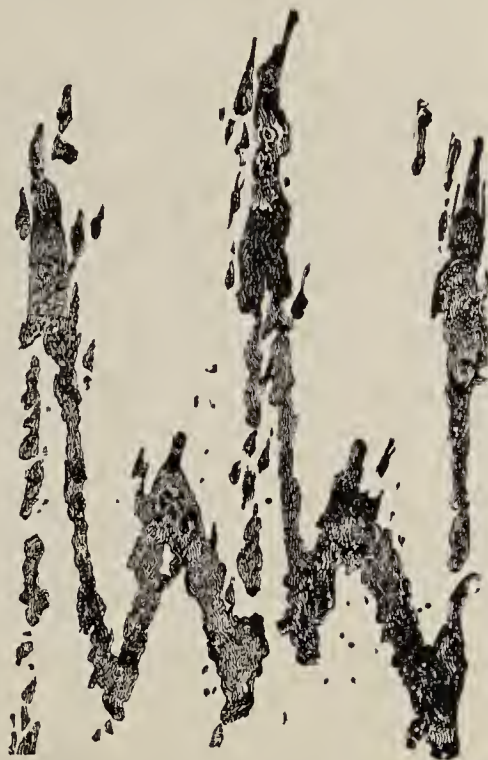


FIG. 43.—HEMAUTOGRAM. (Landois.)

The nature of the dicrotic wave has been much disputed.



Does it, like the percussion wave, travel from the heart towards the periphery ; or is it a wave reflected from the periphery ? It is probably the former, and produced as follows : when the systolic discharge commences, the primary pressure wave starts ; while this discharge continues, the tidal pressure wave is maintained ; when it suddenly ceases, a negative pressure wave starts which gives rise to the dicrotic depression ; the semilunar valves now suddenly close and uphold the column of blood in the aorta ; the cessation of the negative wave and the sudden arrest of the column of blood by the semilunar valves together give rise to a second positive wave starting from the aorta and propagated throughout the arterial ramifications as the dicrotic wave. It is, however, not always easy to identify these so-called tidal and dicrotic waves on all kinds of tracings, nor may we assume that their mechanism is always the same ; recent observations show that *reflected* waves may possibly occur in the arterial system of the dog (v. Frey).

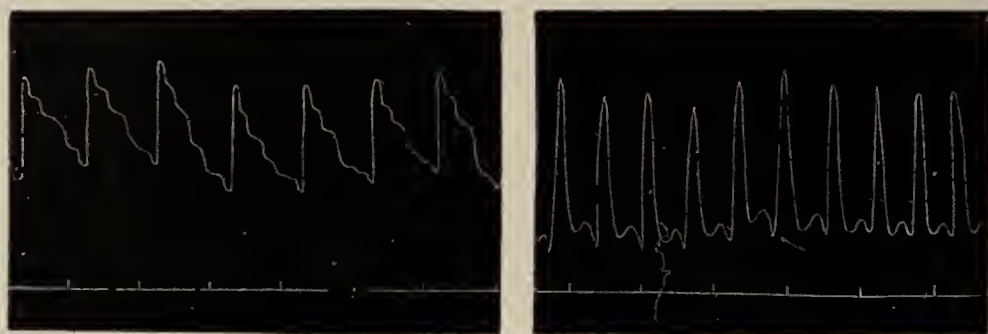


FIG. 44.—PULSE-TRACING BEFORE AND AFTER THE INHALATION OF NITRITE OF AMYL. TENSION DIMINISHED AND FREQUENCY INCREASED.

The sphygmograph has proved useful as an instrument by which certain characters of the pulse have been demonstrated, but it has no valid claim to be regarded as furnishing any clinical information that is not obtained by a skilled touch. A knowledge of sphymographic tracings gives greater definiteness and precision to the sensations obtained by feeling the pulse ; on the other hand, an unguarded use of the instrument, or a too minute interpretation of tracings, to the exclusion of a due regard to the palpable character of the pulse, is very likely to mislead with false evidence. The instrument may, in fact, be an added source of error, and in practice had better be avoided than carelessly used.

*Velocity of the pulse.*—The cardiac impulse and the radial pulse are not absolutely synchronous, as may easily be verified by feeling the two impulses simultaneously—the radial pulse will

be felt a short interval after the cardiac impulse ; or if the heart be listened to while the radial pulse is felt, the latter will be noticed to occur after the first sound, about midway between the first and second sounds. Nor is the pulse simultaneous throughout the arterial system ; for instance, the carotid may be felt to precede the radial pulse, and the femoral to precede the tibial pulse. In short, the cardiac discharge gives a pulse throughout the arterial system propagated with a definite velocity from the heart to points further and further removed. This velocity is measurable by instruments by the method of simultaneous tracings taken on a revolving cylinder. If, for instance, the cardiac impulse and the radial pulse be simultaneously recorded by levers, the points of which are ver-

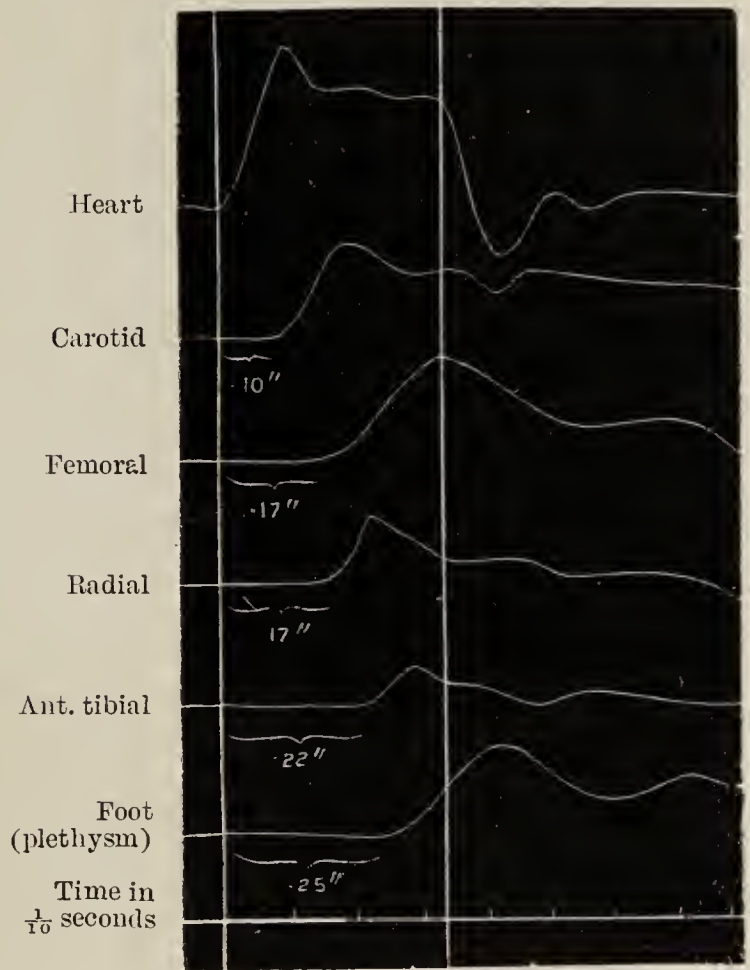


FIG. 45.—THE PULSE-WAVE IN THE ARTERIAL SYSTEM (W.).

tically beneath each other, it will be noticed that the cardiac lever begins to rise before the radial lever, and if the interval between these initial points be compared with a time tracing, as of a vibrating tuning-fork, its time value may be determined with great accuracy. This interval is spoken of as the *pulse-delay* : it is as follows at different points of the arterial system in hundredths of a second.

Heart-carotid interval	.	.	.	.	.	.	10
Heart-radial	„	.	.	.	.	.	17
Heart-femoral	„	.	.	.	.	.	17
Heart-tibial	„	.	.	.	.	.	22
Carotid-radial	„	.	.	.	.	.	7
Femoral-tibial	„	.	.	.	.	.	5

In these measurements it is to be noted—(1) that the heart-carotid delay is disproportionately long ; (2) that the carotid-radial is longer than the femoral-tibial delay. The first of these



points is due to an interval of 6 to 8 hundredths of a second between the commencement of the ventricular contraction and the forcing open of the aortic valves against the resistance of intra-aortic pressure. The second point illustrates the fact that the pulse-wave travels more rapidly in the arteries of the lower than in those of the upper extremity. The average velocity of the pulse-wave is 9 meters per second ; this velocity must not be confused with the velocity of the blood-current, which does not exceed .3 meter per second. The velocity of the pulse-wave is a very constant magnitude ; variations of velocity are comparatively small and not to be detected by feeling the pulse. Clinically, a 'delayed' pulse is usually a weak pulse, or a pulse is erroneously called 'delayed' because it is timed with reference to a diastolic instead of a systolic impulse. The radial pulse never succeeds the second sound of the heart, even when the systole is at its shortest ; normally it is felt to be equidistant between the first and the second sound (w.).

**On the properties and mode of contraction of cardiac muscle.**—The normal heart contracts regularly ; the same heart removed from the body and empty of blood continues for a time to contract regularly—on warm-blooded animals for a few minutes, on cold-blooded animals for several hours. The heart thus differs from all other muscles, which do not contract until they are stimulated, and possesses within itself the conditions of regular and apparently spontaneous action. What are these conditions, and by what portion of the heart's substance are they possessed ? In other words, what is the cause of the heart's rhythm ? Various answers have been given, different theories have been proposed, and the study of the question has brought many facts to light. These we have to consider. It has been answered that the continuance of the rhythmic beat is owing to the presence of ganglia : a theory has been advanced to the effect that these ganglia rhythmically discharge themselves, and thus excite rhythmical contraction of the muscle. This theory was based upon the observation of the fact (?) that on cold-blooded animals, not only the entire organ, but separate bits will continue to beat for a time ; on microscopic examination ganglia were found in such bits, while in other bits which remained motionless ganglia were sought for in vain. Such observations are liable to vary with the expectation of the observer, and they are contradicted by exactly opposite observations ; the lower two-thirds of a frog's ventricle

—a so-called *ventricle-apex* preparation—which contains no ganglia, will, if supplied with nutrient fluid, continue to beat rhythmically; a strip of muscle cut from the ventricle of the tortoise, and destitute of ganglia, may, after a longer or shorter period of quiescence, commence and continue to beat. These last observations are accepted as correct, and ganglia are not now believed to be essential to the continuance of the heart's beat. The property of rhythmic contractility is an attribute of cardiac muscle, and it is useless to ask what the cause of that property is; all that may be said is, that the power of rhythmically beating belongs to cardiac muscle while it is alive, whether in the body or out of the body. Another theory may be mentioned, but only to be dismissed as unfounded. Brücke thought that the rhythmic alternation of action and rest was owing to a rhythmic mode of nutrition; he thought that this was affected by a peculiarity in the blood-supply by the coronary arteries, he imagined that these were shut off from the aorta by its semilunar valves during the systolic discharge, but flushed with blood during the diastole, and he supposed that the diastolic blood-supply and consequent nutrition constitute the cause of each systolic contraction. This theory is quite untenable, it does not account for the action of the excised heart, or of the frog's heart, or of bits of heart; the supposition that the coronary blood-supply occurs during diastole is not even correct—the arterial pulse is systolic in the coronary as in all other arteries. Nor does the heart stop at once after the complete arrest of circulation through the substance of its muscle by ligature of the coronary arteries; it continues to beat for several minutes after this operation, which is indeed equivalent in its effects to the actual excision of the organ.

*Stannius' experiments.*—At a period when the attention of physiologists was focussed upon the ganglia of the heart and their supposed mode of action (1852), Stannius made certain experiments upon the excised frog's heart which have been constantly referred to by subsequent observers and explained in many different ways. Apart from their explanation, which cannot be given as settled, which, indeed, is even more obscure since we have become assured that rhythmic contractility is a property of cardiac muscle *sine* ganglia, the chief facts observed by Stannius were as follows: if a ligature be tightly applied round the heart at the junction of the sinus with the auricle, the auricle and ventricle stand still in diastole while the sinus continues to beat.



If now a second ligature be applied round the heart at the junction of the auricle with the ventricle, the ventricle recommences to beat, quickly at first, but gradually more slowly, while the auricle remains quiescent. The quiescent state of the auricle and ventricle consequent upon the first Stannius ligature closely resembles the state consequent upon vagus stimulation, the arrest being in diastole, and the heart remaining capable of contracting in response to mechanical or electrical stimuli. But that the arrest is not, as might be supposed, due to vagus excitation by the ligature is proved by the fact that the standstill is equally effected by ligature after atropine poisoning, which completely abolishes vagus excitability. The effects of the first and second ligatures have been 'explained' as follows. Ganglia are present (1) in the sinus (Remak's ganglion), (2) in the auricle (v. Bezold's ganglion), (3) at the base of the ventricle (Bidder's ganglion). It has been supposed that 1 and 3 are 'motor,' and 2 'inhibitory'; that the motor influence of 1 and 3 combined is greater than the inhibitory influence of 2, consequently in the absence of all ligature the heart beats; that the motor influence of 3 is less than the inhibitory influence of 2, consequently after the first ligature, cutting off the motor influence of 1, the auricle-ventricle stands still; while after the second ligature, cutting off also the inhibitory influence of 2, the ventricle influenced by 3 alone and unopposed, recommences to beat. Obviously such an 'explanation' explains nothing, and only translates Stannius' statements into other and very doubtful terms without adding anything to our knowledge.

We are, indeed, beginning to modify our statements of the facts themselves. The standstill after the first ligature is not permanent nor even of long duration; half an hour is a comparatively long period for it to last—more usually its duration is only a few minutes, at the expiration of which the auricle-ventricle recommences to beat: sometimes the ligature fails entirely and the auricle-ventricle goes on beating, slowly at first and gradually faster. Again, a ligature in the auriculo-ventricular groove may cause the ventricle of a quiescent auricle-ventricle to resume its beat as above stated, but it may also temporarily arrest the beat of an actively beating auricle-ventricle. It appears rather as if the property of rhythmic contraction possessed by all parts is at its maximum in the sinus, at its minimum in the ventricle, while in the ventricle it is greater at the base than at the apex; contraction normally starts from the point of greatest

instability, *i.e.* from the sinus; removal of the sinus removes a leading part and causes shock, and the beat of the rest of the heart ceases for a period; after a time the beat recommences, contraction starting from the auricle; a second ligature or section removing the auricle may cause a similar result, *viz.* a temporary cessation of the ventricular beat, or if (omitting the ligature at the sinus-auricle junction) a ligature be at once applied to the auriculo-ventricular groove, the ventricle will be temporarily arrested in diastole. In short, though usually the first ligature gives arrest which is ended by the second ligature, yet under varied conditions the first ligature may fail to give arrest, or the second ligature may give arrest. The results are thus not only complex, but variable, and cannot be made to bear any precise significance. No corresponding results have been obtained on the mammalian heart; if all nervous communication between the auricles and ventricles be destroyed, the chambers continue to beat, but with independent rhythms; vagus stimulation now fails to arrest the ventricles, while it still produces auricular standstill.

‘*All or nothing.*’—But if the theoretical significance of Stannius’ experiments is obscure or doubtful, it is otherwise as regards their practical value. The first Stannius ligature affords a constantly employed means of obtaining the frog’s heart in a quiescent state, when it can be excited and employed for experiments that cannot be made while it is spontaneously beating. If a ‘Stanniused’ and consequently quiescent frog’s heart be subjected to stimuli of gradually increasing strength, it is found that the weakest stimulus which produces any effect at all, produces all the effect of which the muscle is capable—in other words, the effect is at once a maximum effect; a single stimulus either produces no effect at all, or it produces the full effect of which the muscle is capable; ‘*all or nothing*’ is the motto of the heart’s contraction under these circumstances. To what extent may we suppose that this holds good for the normal beat? Do a succession of normal systola vary in strength, or are they uniform and maximal? The usual answer is that they are normally maximal and complete, and that the ventricle normally empties itself of its contents completely, but it is obvious that this does not imply equality in successive contractions, for the successive charges vary in amount, being greater or smaller and consequently expelled by greater or smaller contractions; still, the motto ‘*all or nothing*’ holds good; the contraction, whether great or small, is the



greatest possible, *i.e.* maximal, and the normally acting ventricle is completely emptied at each such contraction. Abortive or incomplete contractions are abnormal.

*The staircase.*—A Stanniused heart, excited at regular intervals of, say, two seconds by single induction shocks sufficiently strong to make the heart beat without fail, gives a succession of contractions of increasing magnitude. Each contraction, although

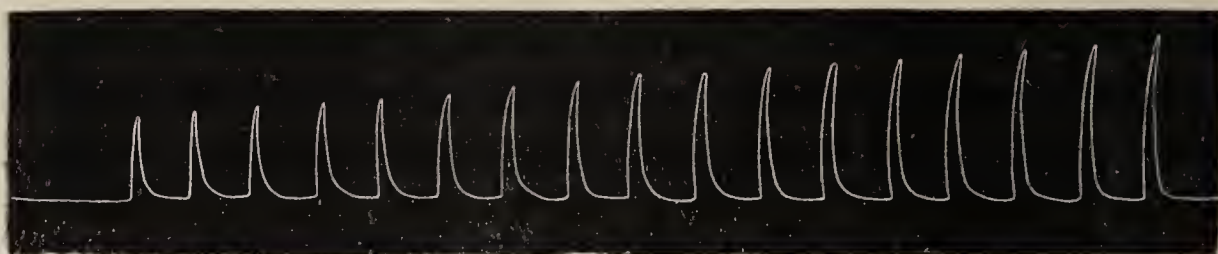


FIG. 46.

‘Staircase’ of beats of a Stanniused frog’s heart excited by maximal induction shocks at intervals of three seconds.

‘maximal,’ inasmuch as it is the full effect of which the muscle is then capable, is a little greater than the preceding contraction, and the general outline of such a series when recorded is that of a staircase, each ascent of the recording lever from the base line reaching a step higher than its predecessor (Bowditch). The effect is not, however, peculiar to cardiac muscle; ordinary voluntary muscle stimulated by a succession of maximal induction shocks usually gives an ascending or staircase group before entering upon the decline due to fatigue.

*The refractory period; tetanus.*—The heart is not equally excitable during rest and during action; it is less excitable during action than during rest; further, it is less excitable during rising action than during declining action, *i.e.* during the beginning than during the end of systole; the comparative inexcitability is so marked during the commencement of systole that this period has been termed the *refractory period*. These variations of excitability are not absolute but relative, the heart is not absolutely inexcitable during systole, but only less excitable than during diastole. Further, the inexcitability is more profound during earlier than during later portions of the systolic period, and a similar statement holds good for the other periods, the state of excitability being at its minimum at first, thence progressively increasing to its maximum, when a second spontaneous beat is about to be discharged (*vide* fig. 47). Similar statements are applicable to excited beats of the Stanniused heart.

Can cardiac muscle be tetanised like voluntary muscle by a rapid succession of stimuli? To this the experimental answer is 'No,' as might be foreseen from consideration of the refractory period. The early stage of each excited contraction is refractory to stimuli, which consequently fail; the refractory state is not absolute, but shorter and shorter to increasing strength of stimulus.

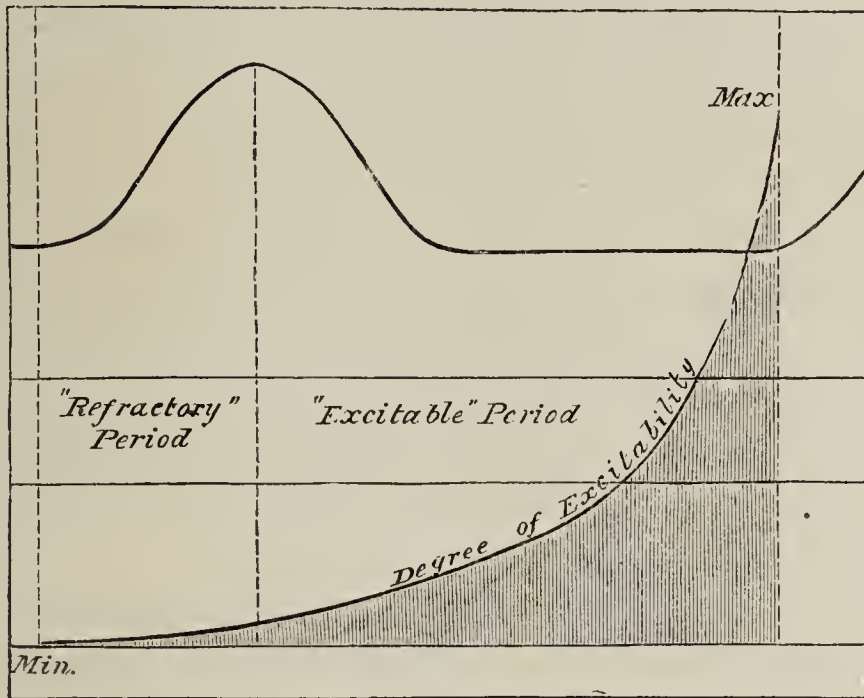


FIG. 47.

To illustrate the varying excitability of a frog's heart at different periods of systole and of diastole. The excitability is lowest during the first half of systole, greatest during the second half of diastole.

Strong, rapid stimuli can consequently increase the rapidity of the beat, and to such an extent as to give an incomplete tetanus, but complete uniform tetanus like that of voluntary muscle cannot be obtained by a succession of stimuli.

*The wave of contraction.*—A contraction, whether spontaneous or excited, is not absolutely simultaneous throughout the whole mass of the ventricle, but it sweeps over the contractile tissue from its seat of origin. This is easily verified by simultaneous tracings of two levers resting on different parts of the ventricle; these do not rise together, but one after the other. A Stanniused and therefore quiescent frog's heart thus treated shows that the wave of contraction can travel in either direction, from apex to base, or from base to apex, according as the ventricle is stimulated at the apex or at the base. A strip of muscle cut from a freshly excised mammalian heart, placed beneath two levers, will contract wave-like from either end to the other, raising one lever after the other, according to the end which is stimulated. This wave-like progress of contraction occurs also in the natural beat; in the frog's ventricle it is from base to apex; the velocity of the wave



can be calculated, *e.g.* if the two levers are 1 cm. apart, and the time interval between the initial points of their elevation  $\frac{1}{10}$  sec., the velocity must be 10 cm. per sec.

In the mammalian ventricle it is far more difficult to obtain evidence of the passage of a wave of contraction, unless the rapidity of movement has been reduced by cooling. In the normal beat of the warm-blooded heart it is impossible to tell by mechanical means whether the contraction begins at the apex or at the base. The only other means of exploration is afforded by electrical instruments; we shall find when we come to speak of these, that the test can be applied to the unexposed as well as to the exposed heart, but we may state now that in the second case we have no assurance that the organ continues to beat normally, while in the first the indications are difficult to decipher with certainty. Any dogmatic statement concerning origin, course, and speed of the wave would therefore be out of place here.

‘*Blocking.*’—If a strip of heart muscle which is still excitable be compressed, and stimulated at one end, the contraction beyond the point of compression is delayed or may be abolished; the passage of the stimulus is more or less *blocked*. If a frog’s heart is set up between the jaws of a clamp which just holds it at the auriculo-ventricular junction, it will while fresh give a series of auricular followed by ventricular contractions. If the clamp be tightened it may give only one ventricular for every two or more auricular contractions. Compression has blocked the passage of stimuli from auricle to ventricle, and instead of every stimulus fulfilling its effect, every other or every third stimulus is alone effectual, in consequence of partial ‘*blocking*’ at the compressed auriculo-ventricular junction (Gaskell).

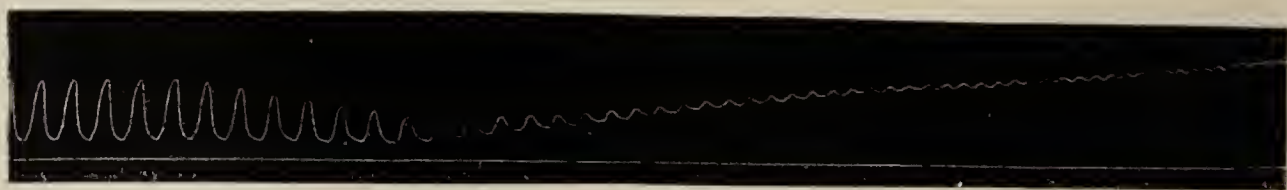


FIG. 48.

Effect of heat upon contraction of frog’s heart. The first effect is diminution of systole; subsequently systolic rigor develops. The frequency of the beats is increased, but in this instance neither ‘heat-standstill’ nor ‘heat-rigor’ is complete.

*Influence of temperature.*—The frequency of the heart’s beat is greater and smaller at higher and lower temperatures respec-

tively, and the duration and extent of each beat are correspondingly smaller or greater. If a spontaneously beating frog's heart be gradually warmed, its beat will gradually diminish in height, and the heart may stop in diastole ('heat-standstill'); on further warming, the organ will enter into the state of tonic contraction known as 'heat-rigor.'

**Drugs by which the action and properties of the heart can be modified.**—Many different drugs modify the heart's action, the effects produced being in certain cases marked and predominant, in others appearing as subordinate features among effects better characterised elsewhere than upon the heart. It is not possible to draw up an accurately-classified list of cardiac drugs under definite headings such as 'cardiac stimulants,' 'cardiac sedatives,' &c.; it is better to take each drug separately and describe its best-characterised effects upon the heart, such as alterations of the spontaneous beats, of the excitability and of the tone of cardiac muscle, or modifications in the excitability of its nerves. We shall find that theories as to the exact parts affected by different drugs are numerous and ingenious, but we shall endeavour to keep these theories very distinct from the tangible facts, and to avoid the description of experimental results in language involving theoretical assumptions. For example, the statement that 'muscarine stops the heart in diastole, atropine sets the heart beating again,' is simply descriptive of evident effects, whereas the statements 'muscarine excites inhibitory ganglia, atropine paralyses them,' or 'muscarine paralyses motor ganglia, atropine excites them,' include suppositions which may or may not be true. The apparently simple proposition, 'muscarine excites inhibitory ganglia,' is in reality composed of the descriptive statement, 'muscarine stops the heart,' and of the more or less probable suppositions that the heart is stopped by inhibitory ganglia, and that muscarine has a specific action upon inhibitory ganglia. We may take this opportunity of formally recognising a distinction which is of universal value, and especially desirable in physiological study. Descriptive statements are definite and open to proof or disproof; inferential statements are liable to be indefinite and impregnable by proof or disproof; composite statements involving in the same words description and interpretation are treacherous and embarrassing. A descriptive statement applies to matter of fact and must always be of intrinsic value, although it may be of more or less importance. An inferential statement conveys explanation of admitted fact, and is sometimes of the



utmost value, but it is dangerous and liable to abuse. A mixed statement is useless or mischievous, and should, whenever possible, be decomposed into its positive and suppositive components. 'Muscarine excites inhibitory ganglia' is a mixed statement. 'Muscarine stops the heart' is a positive statement. 'Inhibitory ganglia stop the heart'—'Muscarine acts upon inhibitory ganglia,' are suppositive statements.

The chief drugs of which the action upon the heart has been most carefully studied in the laboratory are:—*muscarine*, *pilocarpine*, *atropine*, *nicotine*, *physostigmine*, *curare*, *digitaline*, *aconitine*, *veratrine*; *acids*, *alkalies*, and *neutral salts*; *alcohol*, *chloroform*, and *ether*. Of these drugs, *atropine* and *muscarine* are of most interest from a physiological standpoint, while their use as cardiac medicines is very limited; on the other hand, *digitalis*, *aconite*, *ether*, &c., are of far more importance as therapeutic agents, and this has led to their exhaustive examination in the laboratory.

*Muscarine* is an alkaloid extracted from a poisonous mushroom—*amanita muscaria*—present also in putrefying fish or flesh. The effect of a small dose of muscarine injected into the circulation of a dog or rabbit, or directly applied to the exposed heart of a frog, is a rapid diminution of the heart's beat, ending in its complete arrest in diastole. This state closely resembles that of a



FIG. 49.

Effect of Muscarine, and subsequently of Atropine, applied directly to an excised Frog's heart.

heart arrested by vagus stimulation, the muscular substance of the heart remaining responsive to direct stimulation. The generally-received *theory* is that muscarine strongly stimulates inhibitory ganglia of the heart, and that to a less degree it exercises a direct effect upon the cardiac muscle itself, depressing its excitability.

Pickering's observations are to the effect that the embryonic heart (in which a nervous mechanism is not yet developed) is not affected by muscarine.

*Pilocarpine* has an action on the heart similar to that of muscarine.

*Atropine*.—A small dose of atropine injected into the circulation

gives rise to a great increase of pulse-frequency; the effect is similar to that consequent upon section of the vagi, and is correspondingly well marked on dogs, ill-marked upon rabbits. Excitation of the vagus on atropinised animals fails to slow or arrest the heart's action. Thus the effect of atropine is equivalent to an interruption between vagus and heart—a paralysis of the intra-cardiac inhibitory apparatus. On frogs atropine directly applied to the exposed heart gives no appreciable alteration of the beat, but excitation of the vagus or of the sinus-auricle junction fails to arrest the heart—on the contrary, it quickens and augments the beat owing to excitation of accelerator fibres. A heart in diastolic arrest caused by the application of muscarine, recommences to beat shortly after the application of atropine. Atropine is thus *antagonistic* to muscarine.

*Nicotine*, like atropine, abolishes the inhibitory effects of vagus stimulation; stimulation of the vagus of a frog poisoned with nicotine, so far from arresting the heart, actually accelerates it.

*Physostigmine* or *eserine*, one of the alkaloids extracted from the calabar bean, in small doses increases the excitability of the vagus; it is in this respect to a limited degree an antagonist to atropine; a frog atropinised to a point when excitation of the vagus fails to arrest the heart may recover its susceptibility to vagus stimulation after a small dose of physostigmine. After large doses of physostigmine vagus excitation fails to arrest the heart.

*Curare*, which acts more particularly on the termination of nerve in voluntary muscle, may also, if administered to excess, interfere with the action of the vagus on the heart. On fully curarised frog's vagus excitation consequently fails to arrest the heart.

*Digitaline*, the chief alkaloid of *digitalis*, at first exaggerates the excitability of the vagus or actually excites it; the signs of this are, that unusually weak stimuli will arrest the heart, or that the frequency of the beat is actually lowered. As a later effect the excitability of the muscle itself is increased, the sign of this being prolonged and stronger systolic contraction, so prolonged as to merit the name of cardiac contracture. The progressive effects of digitaline upon the heart may be grouped as follows:—*1st stage*, increased strength, increased duration, and diminished frequency of the heart's beat; *2nd stage* irregular, often dicrotic and frequent beat; *3rd stage*, systolic arrest; *post mortem* the heart of frogs is usually found in systole, of dogs



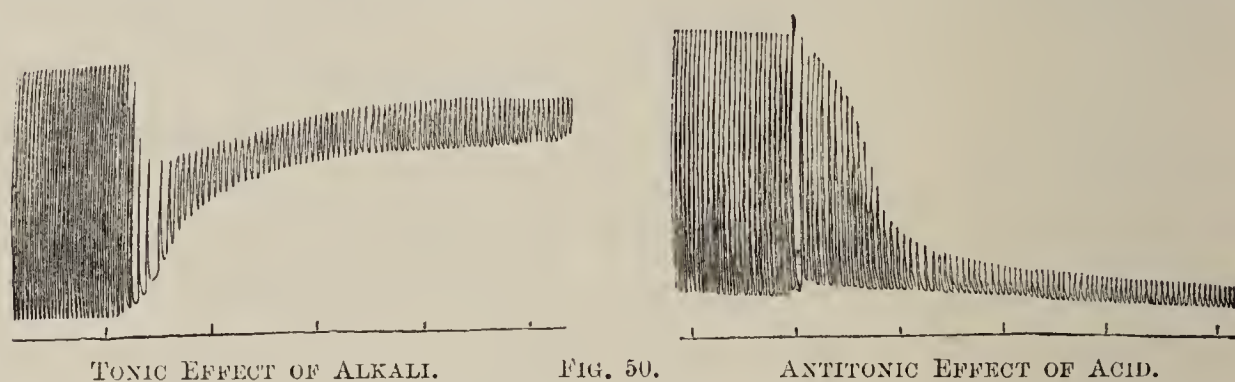
sometimes in systole, sometimes in diastole. Medicinally digitalis is classed as a cardiac tonic.

*Veratrine* has an action on the heart similar to that of digitalis. The prolongation of the contraction is even more marked ; it is abolished by potassium chloride.

*Aconitine*, like digitalis and veratrine, excites the vagus and slows the heart ; on frogs the slowing is preceded by a short stage during which the beat is more frequent ; on dogs and on man this is not the case, the heart's beat becoming slower from the beginning. Medicinally aconitine is classed as a cardiac sedative.

*Alcohol, chloroform, and ether*, in small doses, are cardiac stimulants, causing the heart to beat more rapidly and more strongly. In larger doses they become depressent.

*Acids and alkalies*, in dilute solution, affect the duration of the



Effects of alkali and of acid upon the excised frog's heart. (The recording lever is raised with systole and falls with diastole.)

systole and the tone of the heart. Dilute alkali (caustic potash,  $\frac{1}{20000}$ ) prolongs the duration of systole, and leads to tonic contraction in systole. Dilute acid (lactic acid,  $\frac{1}{20000}$ ) shortens the systole and weakens it, leading to complete relaxation and arrest in diastole, *i.e.* has an antitonic effect. These statements refer to the heart and to the arteries of the frog, through which solutions are made to circulate.

*Neutral salts.*—A solution of sodium chloride in distilled water, .6 per cent., made to circulate through the frog's heart, has no power of maintaining the beats ; these gradually and progressively decline. The addition of a minute trace of calcium chloride restores the beats and causes them to become prolonged. The further addition of a trace of potassium chloride causes these prolonged beats to resume their normal character. Potassium and calcium salts are mutually antagonistic, sodium salts are relatively indifferent. Thus NaCl, CaCl<sub>2</sub>, and KCl in dilute solution compose a fluid which maintains normal beats of the

heart better than the solution of any one, or any two, of these salts alone; such a fluid may therefore be termed a 'nutritive' fluid (Ringer).

'Nutritive' fluids.—The nutritive quality of such a fluid may be improved by the addition of a minute trace of serum-albumin, milk, blood, or blood-serum. When any of these fluids are made to circulate through the frog's heart, it may continue to beat normally for many hours, and the modifying influence of drugs can be conveniently observed by adding them to the nutritive fluid. Egg-albumin, albumose, and peptone have no nutritive action upon the heart, or even a contrary effect; the only proteid possessing a true nutritive action is serum-albumin (Kronecker).

*Perfusion-cannula and frog-heart apparatus.*—Many of the results enumerated above, in so far as they relate to the behaviour of the frog's heart, have been obtained by means of an apparatus permitting a record of the contractions to be taken while solutions of different substances are made to flow through the heart. The frog-heart apparatus in most general use are those of Roy and of Kronecker; in both the essential portion is the two-way or perfusion-cannula, which is introduced through the sinus venosus and auricles into the ventricle, and tied in so as to serve as the sole inlet or outlet; in Kronecker's apparatus the movements of the heart are recorded by a mercurial manometer; in Roy's, the heart works in a vessel full of oil fitted up on the principle of the oncograph. To test the nutritive action of a fluid, the proceeding is as follows:—The heart is first to be 'washed out' by letting a stream of normal saline flow through the perfusion-cannula and heart until the spontaneous beats and the electrical excitability have disappeared; the test solution is now allowed to flow through the heart, and is known to be 'nutritive' or not accordingly as electrical excitability and spontaneous contractions are or are not restored.

**Physiological anatomy of cardiac nerves.** The cardiac nerves are formed by fibres derived from (1) the vagus, (2) the sympathetic. Both sets of fibres are efferent in function; the former restraining, the latter stimulating, the activity of the heart. A third set of fibres has been experimentally demonstrated in mammalia (cat, dog, rabbit), afferent in function named *depressor*, on account of their influence upon the blood-pressure, and taking a somewhat variable course in the vagus or in its branches.

*Frog.*—The vagus in the frog is in reality composed of two nerves—



(1) the vagus proper, which is purely *inhibitory*, (2) a sympathetic branch, which is purely *augmentor* of cardiac activity. The vagus proper has its origin or centre in the medulla oblongata, and after emerging from the jugular foramen is joined by a branch of the sympathetic

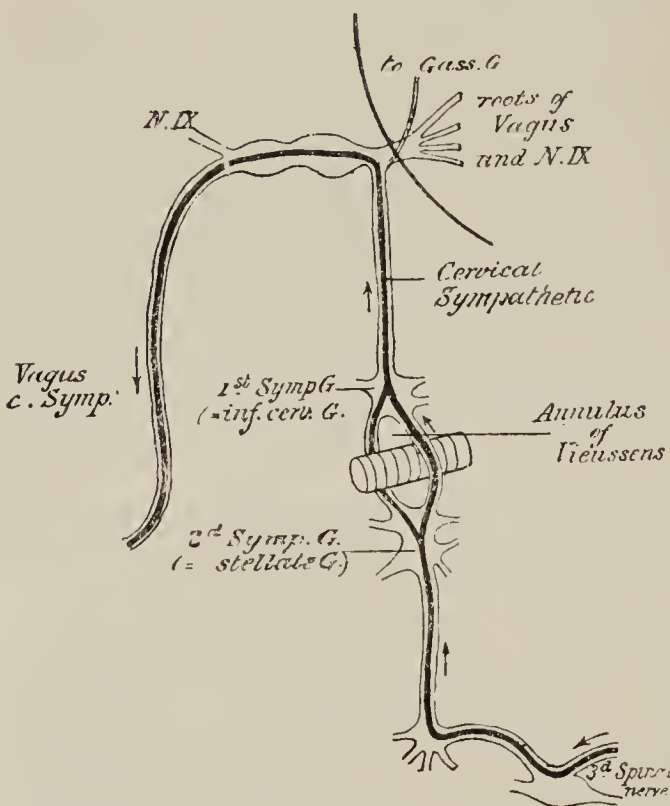


FIG. 51.—CARDIAC NERVES OF FROG. (Foster.)

derived from the spinal cord by way of a communicating branch from the third spinal nerve to certain sympathetic ganglia which may be regarded as analogous with the inferior cervical and first thoracic ganglia of mammalia (Gaskell). The conjoint vago-accelerans passes along the superior vena cava to the sinus venosus, thence along the auricular septum to the auriculo-ventricular ganglia.

*Mammalia.*—The cardiac nerves of mammalia (cat, dog, rabbit) comprise (1) inhibitory fibres to the heart, (2) accelerator fibres to the heart, (3) depressor fibres from the heart. The *in-*

*hibitory* fibres are derived from the spinal bulb by the internal branch of the spinal accessory; they run downwards in the vagus nerve, and reach the heart by its cardiac branches. The *accelerator* fibres have their origin in the spinal cord; leaving it by the anterior roots of the second and third thoracic nerves, they reach the sympathetic by the white rami communicantes of these two nerves, and pass upwards through the first thoracic or stellate ganglion, annulus of Vieussens, and inferior cervical ganglion, from which they reach the heart through the cardiac nerves in company with the cardiac fibres of the vagus. The *depressor* fibres have their origin in the heart, pass along its cardiac nerves or (in the cat and rabbit) as a definite nerve which can be anatomically as well as experimentally isolated, join the superior laryngeal and vagus, and thus reach the spinal bulb. As regards the minute anatomy of these fibres, it has been shown by Gaskell that the cardio-inhibitory fibres are small medullated fibres throughout their course in the spinal accessory, vagus and cardiac nerves, and probably remain medullated down to the cardiac ganglia. Cardio-accelerator fibres are small medullated fibres in the anterior roots, rami communicantes, and sympathetic channels up to the stellate and inferior cervical ganglia. From this point onwards in the cardiac nerves they are non-medullated.

**Physiological anatomy of vasomotor nerves.**—The vaso-constrictor nerves of the whole body leave the spinal cord by the anterior

roots of the thoracic nerves, and enter the sympathetic system by white rami communicantes—in the dog by the sixteen pairs of nerves from the second thoracic to the fourth lumbar (Gaskell). The rami communicantes of the upper thoracic nerves pass to the cervical ganglia and form the cervical sympathetic; they also furnish the vaso-constrictor nerves of the anterior extremities by filaments that pass from the stellate ganglion to the brachial plexus. The lower thoracic nerves give off rami communicantes which form part of the abdominal splanchnic nerves, and furnish the vaso-constrictor nerves of the posterior extremities. The course of vaso-dilatator fibres is far more doubtful, owing to the fact that their presence at any given point is difficult of demonstration; all we may say with certainty is that, like vaso-constrictors, they take origin from the spinal cord and emerge by anterior nerve-roots. It is probable that dilating may be as widely distributed as constricting nerves, but their actually ascertained existence is practically limited to nerves of the head and of the pelvis (chorda tympani and nervi erigentes); and although we shall have occasion to refer to vaso-dilatation caused by the excitation of the sciatic and of the splanchnic nerves, we shall find that the facts themselves are not sufficiently assured to justify anatomical descriptions. According to Gaskell, however, vasomotor

nerves are microscopically distinguishable from spinal nerves, and the two kinds of vasomotor nerves are distinguishable from each other; both kinds of vasomotor nerve-fibres are of the small medullated kind (from  $1.8$  to  $3.6\mu$  in diameter) and leave the cord in the anterior nerve-roots; vaso-constrictors pass from the spinal nerves to the sympathetic chain by white rami communicantes, and, having lost their medullary sheath in a proximal series of ganglia (*i.e.* those composing the main sympathetic chain in the thorax and abdomen), return to the mixed nerve

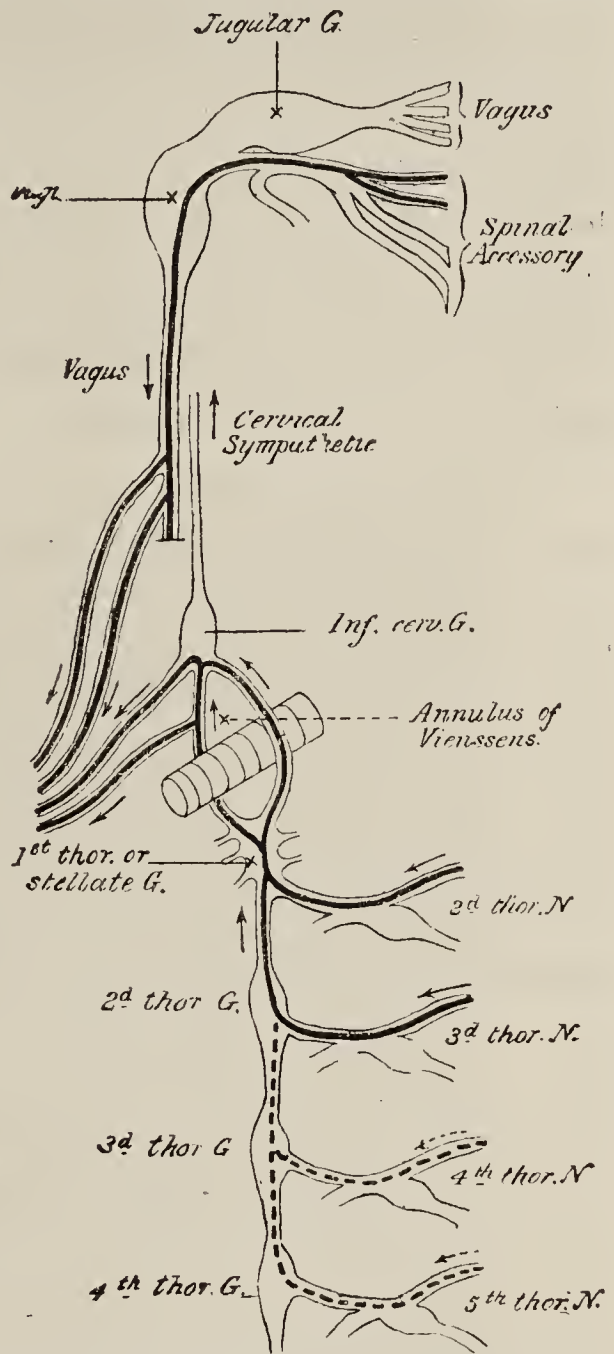


FIG. 52.—CARDIAC NERVES OF DOG. (Foster.)



by grey rami communicantes, vaso-dilatators keep in company with the spinal nerves and remain medullated until they reach a distal series of ganglia (*i.e.* the prevertebral lumbar and sacral ganglia in the pelvis and the submaxillary ganglion on the chorda tympani). Thus from an anatomical as well as from a physiological point of view cardiac and arterial nerves may be classed in two groups, (1) vaso-constrictor and cardio-accelerator, (2) vaso-dilatator and cardio-inhibitory. Nerves of the first group, composed of fine non-medullated fibres, excite muscular action; nerves of the second group, composed of fine medullated fibres, restrain muscular action. On speculative grounds nerves of the first group have been characterised as 'katabolic,' nerves of the second group as 'trophic' or 'anabolic.' The only definite evidence at present offered in favour of this view is an experiment by Gaskell, showing that under certain special conditions vagus excitation gives a positive electrical change in cardiac tissue, *i.e.* the reverse of what happens in ordinary muscle of which the motor ('katabolic') nerve is excited (p. 389).

*Physiological action of cardiac and vascular nerves.*—The heart can beat independently of the central nervous system, but it is nevertheless subject to and controlled by nervous influences transmitted from the spinal bulb and cord by the vagus and sympathetic nerves. Similarly, the vessels—especially the small arteries—are subject to nervous influences transmitted from the bulb and cord by the sympathetic nerves. In each case these influences may either stimulate or restrain the contraction of the heart and of the arteries. To the heart the vagus or pneumogastric nerve is the channel of restraining or inhibitory influence, the accelerator nerves are the channels of stimulating or accelerator influence. To the vessels the sympathetic nerves are channels of influences which increase as well as of influences which diminish their contraction—of constricting as well as of dilating influences. It is by these nerves that the circulation is regulated; the cardiac nerves modify the beat of the heart, the vasomotor nerves modify the state of the peripheral arterioles, causing the general and local alterations of blood-pressure and of blood-flow which have been described above (p. 59).

**The vagus: effect of division.**—If one vagus be exposed and divided—on a dog, for instance—the heart's action undergoes little change; if anything, its beat may be somewhat more frequent than before the operation. If the other vagus be divided, the frequency of the heart-beat will be much increased—in some cases up to twice or three times that observed before the operation (on the dog; the effect is less marked on the rabbit). These

facts show that the vagi are channels of a constantly exerted restraining or inhibitory influence which checks the action of the heart; when this restraint is abolished by section of both vagi, the heart beats on with greater frequency.

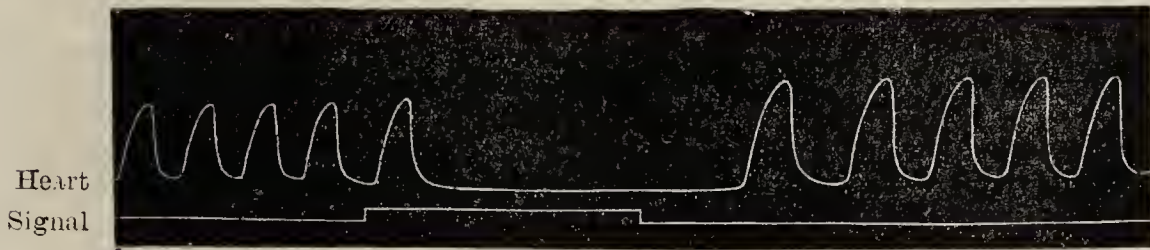


FIG. 53.

Effect of vagus excitation upon contractions of frog's heart; the time of excitation is indicated by the rise and fall of the signal line.

*Effects of excitation.*—If the peripheral end of one of the divided vagi be stimulated by induction currents, the frequency of the beat will be reduced to or below the normal according to the strength of stimulation; or, if the latter be sufficiently

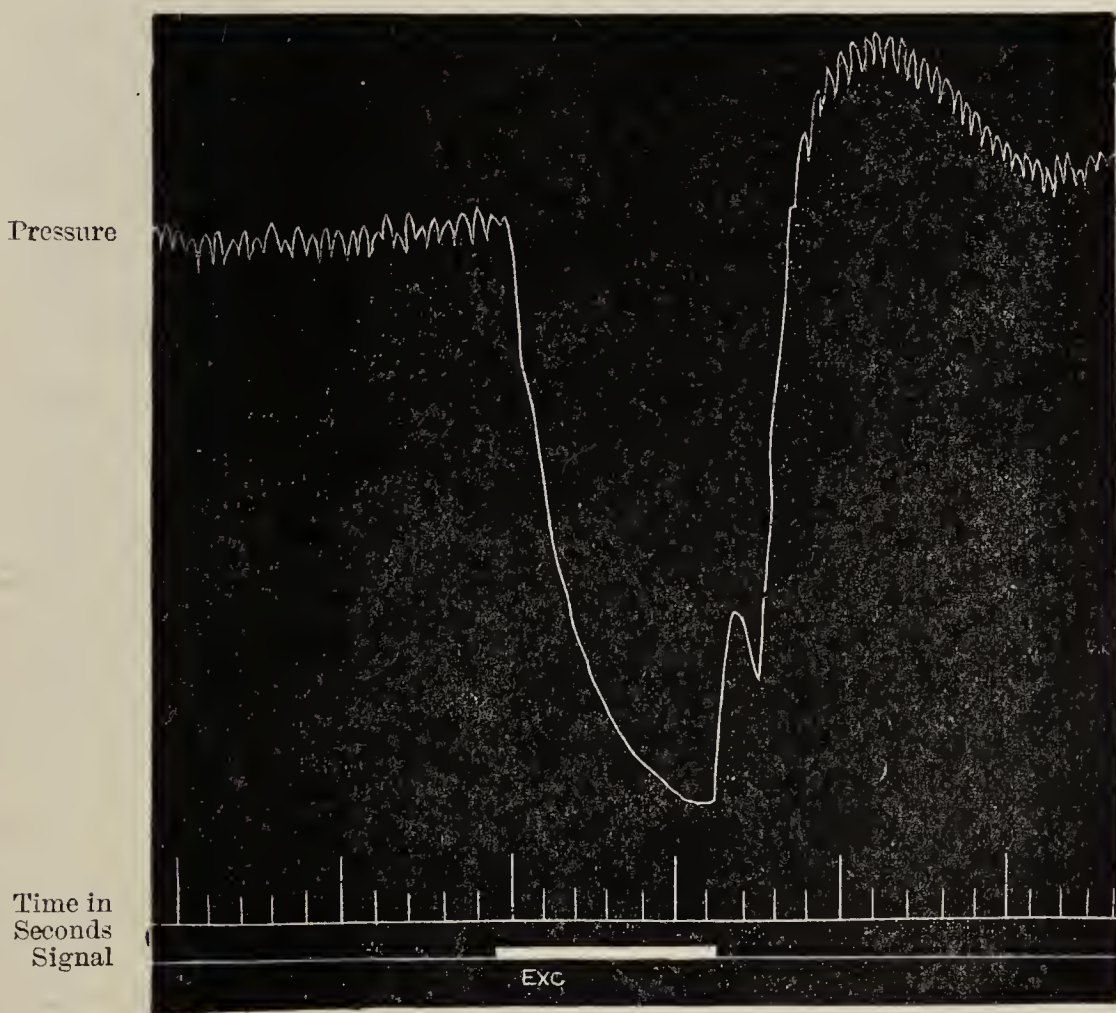


FIG. 54.

Blood-pressure tracing from a rabbit. Excitation of the peripheral end of the right vagus; fall of blood-pressure due to arrest of heart.

strong, the beat will be prevented altogether, and the heart will 'stand still' in a state of diastole. Division of the vagi lets the heart go, their stimulation holds the heart in. The vagus has



thus a precisely opposite action to that possessed by the motor nerves of muscles; it does not excite but restrains cardiac contraction—it is an *inhibitory* nerve (Weber, 1846).

Does the vagus communicate directly with the cardiac muscle, or only indirectly by the intermediation of ganglia? Or, to put the question otherwise, Does the vagus directly interfere with the action of the muscle, or does it interfere with the motor action of peripheral ganglionic organs? No assured answer can be given to the question. Upon the known fact that the action of the vagus diminishes the frequency and force of the beat, the opinion has been based that the vagus terminates in the muscles as well as in the ganglia, diminution of frequency being regarded as the sign of vagus action upon the ganglia, diminution of force as the sign of vagus action upon the muscle.

The effect of vagus stimulation is not produced immediately, but a short period elapses between the moment of stimulation and the consequent arrest. This *latent period*, which it is obviously impossible to estimate exactly (Donders roughly estimated it at .05 second on the rabbit), is usually of such length that at least one heart-beat occurs after the commencement of stimulation.

The effect of vagus stimulation, applied for a short period, does not cease immediately with cessation of stimulation, but outlasts it, and is then termed the '*after-effect*'; the most typical vagus after-effect is a continuation of arrest followed by gradual recovery to and beyond the normal. Prolonged stimulation of the vagus is incapable of keeping the heart arrested for an indefinite period, although normally the vagus is in constant untiring action; the heart recommences to beat during the vagus stimulation, and may even beat more strongly than before.

If stimulation of one vagus is kept up until the vagus arrest has come to an end, stimulation of the second vagus will stop the heart. It must, however, be borne in mind that the two vagi are not always equal in their power over the heart; the vagus of the right side is commonly more effectual than that of the left side, though sometimes the case is reversed, and sometimes there is no observable difference.

The short duration of the effects of vagus stimulation is perhaps in part due to the comparative coarseness of experimental stimulation; but it is, doubtless, also due to the presence

in the vagus nerve of accelerator fibres which come into action as the action of the true vagus wears off. Such accelerator fibres have recently been anatomically defined in the frog's vagus. Gaskell has isolated and stimulated, on the one hand accelerator filaments, on the other hand the vagus *minus* such accelerator filaments, obtaining in the first place acceleration, in the second case prolonged arrest of the heart—far more prolonged than can be obtained by stimulating the intact mixed vagus. Accelerator fibres have not been anatomically defined in the mammalian vagus, but experimentally their existence is, if not proved, made highly probable. The *typical effects* of moderate vagus stimulation are diminished force and diminished frequency of the beats of auricle as well as of ventricle, followed by increased force and increased frequency; *exceptionally*, there is no primary diminution, but an immediate increase of force and of frequency. It is therefore probable that the vagus, or properly speaking the vago-sympathetic, in mammals as well as in amphibia, contains at least two kinds of cardiac fibres—inhibitors, which give the typical effects, and accelerators, which give the exceptional effects. Excitation of a fresh vagus with a vigorous heart gives the typical inhibitory effects; excitation of a fatigued or otherwise depressed vagus with a sluggish heart is more apt to give the exceptional effects—primary augmentation and acceleration of the beat. Direct faradisation of the normally beating heart usually gives inhibition: direct faradisation of a moribund heart, which has almost or just ceased to beat, temporarily restores the rhythmic beat.

Vagus inhibition affects both auricle and ventricle, but the former more effectually and profoundly than the latter; it often happens that only the auricle is arrested, the ventricle continuing to beat on its own account, and if while both chambers are in vagus arrest their mechanical excitability be tested, that of the auricle will be found to be far more profoundly depressed than that of the ventricle.

Within certain limits the action of cardiac muscle is more or less easily modified by nervous influences (augmentor as well as inhibitory) according as the temperature is higher or lower.

On mammalia the inhibitory fibres of the vagus are derived from the *spinal accessory* by its internal branch; if the spinal accessory be torn out by the roots and its fibres allowed to degenerate, excitation of the vagus will no longer produce cardiac inhibi-



tion (Waller, 1856). The inhibitory effects of the vagus can be brought about *on man* by mechanical compression of the nerve; on criminals they have been obtained by electrical excitation of the vagus immediately after decapitation. The vagus continues to possess influence over the heart (both auricle and ventricle) of mammalia for as long as the organ continues to beat after death; whether its action be slow and regular, or rapid and tumultuous as sometimes occurs, vagus stimulation continues effective and causes an immediate standstill. This fact affords a convenient means of demonstrating the 'quelling' action of the vagus upon the movements of the heart.

Arrest of the heart may be also brought about by a reflex action of the vagus; mechanical stimulation of the intestine of the frog, or of the tail of the eel, stimulation of the central end of most afferent nerves, including the vagus itself, will cause this reflex inhibition; it is not produced after section of the vagi, nor after destruction of the spinal bulb, thus proving that the vagi are really the efferent channels of the arrest produced by peripheral stimulation. *Fainting* is in many cases due to a temporary arrest of the heart by reflex inhibition. The variations in the heart's frequency which accompany respiratory variations of blood-pressure disappear after section of the vagi; the *expiratory diminution* of frequency (*vide* p. 143) is therefore due to vagus action, and is probably a reflex caused by some form

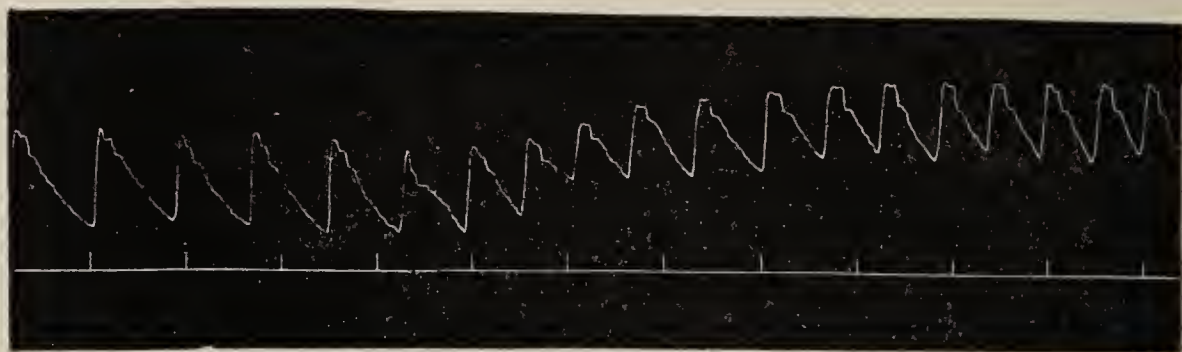


FIG. 55.

Effect of sipping water upon pulse-frequency; the heart beats faster owing to diminished vagus control, and the blood-pressure is raised (man).

of pulmonary stimulation. The act of *swallowing* temporarily interferes with the constant inhibitory action of the vagus on man; sipping a glass of water may thus cause the heart's frequency to rise twenty or thirty beats per minute; in the dog, however, the heart's frequency and the blood-pressure are lowered during deglutition.

*Accelerator nerves.*—The converse of inhibition, as regards the heart, is acceleration or augmentation ; accelerator influences are conveyed from the spinal cord to the heart by the channel of branches of the sympathetic which form part of the cardiac nerves and plexus, their exact anatomy differing in different animals. Section of these nerves is without any constant result as regards the heart's beat ; they are not continuously but only *occasionally* conveying accelerator influence, unlike the vagus, which is *continuously* conveying inhibitory influence. On the other hand, the *excitation* of the peripheral ends of divided accelerators has a uniform and well-marked effect—the frequency and the force of the heart's beat are increased. Compared with the vagus effects as regards latency and duration, the accelerans effect is slowly produced and slowly develops to its maximum. The inhibitory and accelerator nerves are not regarded as true antagonists ; if both be simultaneously excited to action, the effects do not balance and neutralise each other, but arrest during stimulation and acceleration after stimulation are the usual consequences. It is by these channels that accelerator influences reach the heart when the spinal cord is directly stimulated.

*Vasomotor nerves.*—The arteries are not inert pipes, but essentially muscular tubes that are played upon and excited to contract or to dilate by vasomotor nerves ; nerves, the excitation of which causes contraction of the arterial muscle, are called *vaso-constrictors* ; nerves, the excitation of which causes dilatation, are called *vaso-dilatators*. It is easy to understand the action of a constricting nerve, it is difficult to understand that of a dilating nerve ; it is certain, however, that vaso-dilatation does take place as an active, positive, and immediate change, and not merely as a negative and passive change, or as an after-effect, a mere diminution of previous vaso-constriction. The action of a dilating nerve is closely analogous with that of the cardiac inhibitory nerve—the vagus. Each interferes with and suspends the action of muscle—of cardiac muscle in the one case, or of arterial muscle in the other. Vaso-constrictors are widely distributed throughout the body ; their existence was first discovered in the cervical sympathetic, and has since been demonstrated in the splanchnic nerves and in the nerves of the extremities. Vaso-dilatators are also widely distributed throughout the body, though in consequence of the preponderating influence of vaso-constrictor fibres their action is in many cases masked. Their existence was first discovered on the chorda



tympani, as regards its action upon the vessels of the submaxillary gland; they have since been demonstrated in the nervi erigentes, in several branches of the trigeminal, in the nerves of the extremities, in the purely muscular nerves, and even in the cervical sympathetic itself.

Among the long series of experiments alluded to above, two in particular are fundamental to the subject of vasomotor action; these are (1) the experiment of the cervical sympathetic on the rabbit (B. Séquard, Bernard, 1852, Waller, 1853), (2) the experiment of the chorda tympani on the dog (Bernard, 1858).

*Cervical sympathetic of rabbit.*—If the cervical sympathetic of one side be exposed and *divided*, the vessels of the corresponding side of the head become dilated and the temperature of the parts is increased. The division of the nerve has interrupted constant vaso-constrictor influence passing upwards in the cervical sympathetic. If now the cephalic end of the nerve be *excited*, the previously-dilated vessels contract and the temperature of the parts is diminished. This double experiment, the results of which are clear and uniform, marks out the cervical sympathetic as the type and representative of vaso-constrictor nerves, which, as we shall learn from other experiments, are widely distributed throughout the body.

*Chorda tympani of dog.*—Section of the chorda tympani does not produce any definite or constant effects upon the circulation of the submaxillary gland. Excitation of the peripheral end of the nerve causes an immediate and striking change. The previously pale gland becomes vividly red, and whereas previous to excitation of the nerve black venous blood dribbled out of a divided vein, the blood is now bright scarlet and escapes in jets. All these changes prove that the small arteries of the gland have become dilated, so that the arterial blood passes rapidly and with little change through the capillaries, and the arterial pulse is propagated through them into the veins, giving a venous pulse.

Experiments upon the *nerves of the limbs*—more particularly upon the sciatic—have been numerous and their results conflicting. Diametrically opposite assertions have been made—that the sciatic contains only constricting fibres—that it contains only dilating fibres—that it contains both kinds of fibres. The last assertion is probably correct, and the conflicting data are presumably due to differences in the methods and in the circumstances of experiment. The usual effect of section of the

sciatic nerve is vaso-dilatation, which however gradually subsides, the paralysed vessels recovering their original 'tone' in the course of a few days or weeks; the usual *immediate* effect of excitation of the peripheral end of the divided nerve is a brief vaso-constriction, which however soon gives way to vaso-dilatation. As regards methods of observation, the following tests have been employed:—temperature observations, manometer observations, changes in colour and changes of a bleeding surface; vaso-constriction is indicated by a fall of temperature, by pallor of the skin, by a fall of venous pressure or by less copious bleeding from a wound; vaso-dilatation by a rise of temperature, by flushing of the skin, by a rise of venous pressure or by more copious bleeding from a wound. The temperature test gives less immediate information than the others; the thermometer is a comparatively sluggish instrument reacting slowly to changes of temperature, moreover changes of temperature are not instantly produced by changes of the circulation, so that it might easily happen that brief changes of the circulation should escape notice by this method. And, indeed, it is to be remarked that the thermometer has been the instrument most relied on by those observers who have denied the existence of vaso-constrictors in the sciatic nerve. As regards the circumstances of experiment, those most influencing the results are the state of the nerve, the surrounding temperature, and the frequency of stimulation. Excitation of a fatigued nerve, or of a nerve which in consequence of previous section has begun to degenerate, is more apt to give signs of vaso-dilatation, while excitation of a fresh nerve usually gives signs of vaso-constriction; it appears as if vaso-constrictor were less resistant than vaso-dilatator fibres, the former being the first to give way when the nerve is fatigued or degenerating. If the surrounding temperature is high, excitation of the nerve is more apt to cause vaso-constriction than if the surrounding temperature is low, when vaso-dilatation is more likely to be produced; it appears as if in each case the state of the vessels became reversed by the nerve excitation; in the warm medium they are dilated and excitation constricts them, in the cold they are contracted and excitation dilates them. Rapidly repeated stimuli are more suitable to the demonstration of vaso-constriction, a succession of single stimuli at longer intervals (1 to 2 sec.) to that of vaso-dilatation.

*The splanchnic nerves* govern a very large vascular district—that of the digestive viscera. The effects of their division and of



their excitation are of corresponding magnitude, and such as to cause great variations of the general blood-pressure. Physiologically they are vaso-constrictors, as is unmistakably shown by section and excitation of their peripheral cut ends. Section causes paralysis of the intestinal vessels, which dilate and accommodate a large quantity of blood; this is derived from the remainder of the system, and the general blood-pressure consequently falls—to  $\frac{1}{2}$  or it may be to  $\frac{1}{3}$  of its original height—to such a degree in fact that death is imminent or actually occurs owing to the deficient circulation. Excitation of the cut nerves temporarily restores the blood-pressure to or above its original height, in consequence of constriction of the mesenteric and portal vessels. These efferent nerves are in intimate functional relation with an afferent nerve, the depressor nerve, the excitation of which causes reflex inhibition of splanchnic vaso-constriction.

*The nervi erigentes* arise from the second and third sacral nerves, and, passing through the hypogastric plexus, are distributed to the penis, bladder, and rectum. Section of these nerves causes no obvious change, but excitation of the peripheral end causes erection of the penis attributable to vascular dilatation (Eckhardt). Excitation of the N. pudendus has an opposite effect, from which we must conclude that it contains vaso-constrictor fibres (Lovèn).

*The dog's mouth.*—In consequence of the striking results on the rabbit's ear of excitation of the cervical sympathetic, we are led to consider this nerve as the typical example of vaso-constrictor nerves. Schiff in 1856 came to the conclusion that it is also a dilating nerve—a view that is supported by the more recent and no less striking experiment by Dastre and Morat on the dog. Excitation of the upper end of the divided nerve causes the corresponding half of the mouth (tongue, lips, cheeks, and gum) to flush immediately and vividly.

The existence of *pulmonary vasomotor* nerves has been positively affirmed and as positively denied; they probably do not exist (w), or, if they exist, their action is quite insignificant. Cerebral vasomotor nerves have been diligently sought for but not substantiated by experiment.

**Central control.**—The regulation of the vascular system is administered by the central nervous system, viz. the medulla oblongata or spinal bulb, and the spinal cord, from which the vascular and cardiac nerves take origin, the particular parts from

which they spring being spoken of as their 'centres'<sup>1</sup> and comprising (1) the vagus centre in the bulb, (2) the accelerator centre in the cord, (3) the principal vasomotor centre in the bulb, (4) accessory vasomotor centres in the cord. The mode of action of these centres has been experimentally examined by observing (a) the consequences of their destruction or direct stimulation, (b) the consequences of stimulation of afferent nerves before and after destruction of the bulb or cord.

*Destruction of the spinal bulb* abolishes, among other actions, that of the vagus and that of the vasomotor nerves—*i.e.* with its destruction of the vagus and the vasomotor centres are destroyed. In consequence of the destruction of the vagus centre, (1) the heart beats more rapidly; (2) excitation of any afferent nerve fails to produce reflex inhibition of the beat. In consequence of the destruction of the vasomotor centre, (1) the small arteries throughout the body are relaxed, and arterial blood-pressure is lowered; (2) excitation of any afferent nerve fails to produce constriction of the arteries and reflex rise of blood-pressure. The fact that destruction of the bulb entails greater frequency of the heart's beat and fall of blood-pressure, shows that the vagus and vasomotor centres are normally in constant action maintaining the heart inhibited, and the arterioles contracted.

*Excitation of the spinal bulb* provokes, among other actions, that of the vagus and that of the vasomotor nerves. In consequence of excitation of the vagus centre, the heart beats more slowly or is arrested. In consequence of excitation of the vasomotor centre, the small arteries throughout the body are contracted and the arterial blood-pressure is raised. Excitation of the bulb producing these effects may be caused in several different ways—1, by direct electrical stimulation of the bulb itself; 2, by stimulation of the bulb by venous blood; 3, by reflex action of the bulb in consequence of stimulation of afferent nerves; moreover, the vasomotor centre may vary in its actions spontaneously—*i.e.* without assignable cause; and in the moribund state when respiration has already ceased, it may rhythmically wax and wane in action, before it ceases to act altogether (p. 144).

<sup>1</sup> Anatomically a centre is a nucleus or ganglion of grey matter from which nerves originate; histologically this grey matter is composed of nerve-cells with which nerve-fibres are connected. Physiologically a centre is a mediator between centripetal and centrifugal nerve impulses; it receives centripetal impulses from afferent nerves, and emits centrifugal impulses by efferent nerves. 'Spinal bulb' and 'medulla oblongata' are used indifferently as synonymous terms.



From what has been said it will be clear that of the two possible vasomotor changes—constriction and dilatation—the former is the dominant and more marked change; if all the vasomotor nerves of the body are stimulated, as by stimulation of the spinal bulb, the resultant is vaso-constriction and rise of blood-pressure; conversely, if all the vasomotor nerves are put out of action, as by destruction of the bulb, the resultant is abolished vaso-constriction and fall of blood-pressure. As regards reflex vasomotor changes, the effects differ with the afferent nerve excited, and with the state of the vessels. The usual result of stimulation of an afferent nerve is a reflex rise of blood-pressure, but in the case of one particular nerve which is an

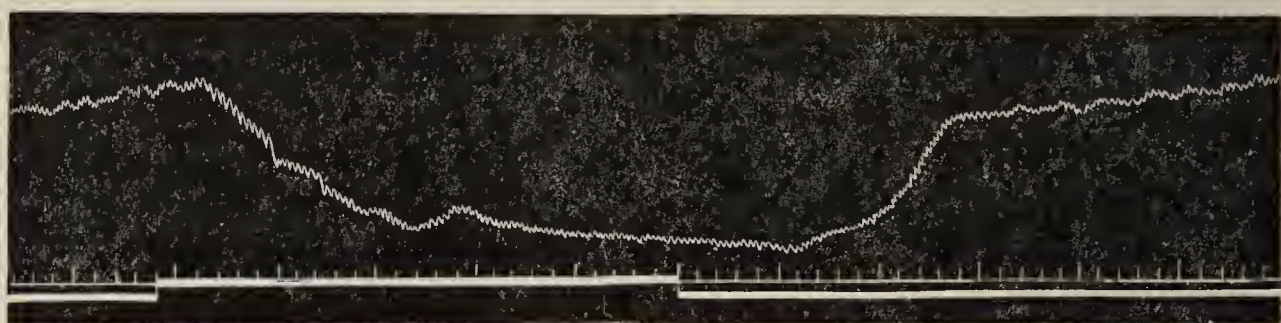


FIG. 56.

Excitation of the central end of the depressor nerve of rabbit. Fall of blood-pressure due to reflex relaxation of the splanchnic vessels. (N.B. The vagus was uncut, hence the fall is associated with reflex slowing of the heart-beat. The abscissa has been raised 3 cm.)

afferent channel from the heart to the bulb—*i.e.* the *depressor*—stimulation of the central end of the divided nerve gives a reflex fall of blood-pressure; the mechanism of this fall is peculiar—it is far less marked if the splanchnics have been previously divided, the effect of their division being of itself a considerable fall of blood-pressure by dilatation of the intestinal vessels. It appears, therefore, that the depressor is not in constant action and that it cannot always be brought into action by stimulation, but that a high blood-pressure is a necessary condition. And it is probable that normally it is brought into action when blood-pressure is so high as to embarrass the heart's action, the depressor then conveying from the heart impulses which depress the vasomotor centre, especially as regards its constant constrictor effect upon the intestinal vessels by way of the splanchnic nerves: these vessels being thereby relaxed, pressure falls, and the heart is relieved. This, indeed, is not the only way in which a high blood-pressure brings about its own remedy; high pressure gives rise to a slower heart-beat (by vagus-control), and with that

slower beat, although the output at each beat is greater, the total output in a given time is less. We may now fully appreciate how the cardiac and arterial factors of blood-pressure are co-ordinated through nervous channels, effecting the physiological combinations alluded to above, viz. greater action of heart linked with vaso-dilatation, and smaller action of heart with vaso-constriction. A greater pulse-frequency has as its mechanical effect a greater blood-pressure; a greater blood-pressure has, as its physiological consequences, a reduction of blood-pressure (*via* depressor and splanchnic) and a reduction of pulse-frequency (*via* the vagus nerve). Guided by these considerations, we may realise that a raised blood-pressure with a greater pulse-frequency, or a lowered blood-pressure with a smaller pulse-frequency, must be of cardiac origin; while a raised blood-pressure with a smaller pulse-frequency, or a lowered blood-pressure with a greater pulse-frequency, must be of arterial origin. The first two of these four cases are illustrated by vagus action, the last two by the effect of dyspnœa and of amyl nitrite.

Stimulation of the central end of any other afferent nerve usually gives reflex rise of pressure, but not always. Repeated stimulation produces each time less and less rise, until, finally, it may produce a fall. It would appear that vaso-constrictor action, by reflex as well as by direct experimental stimulation, wears out more rapidly than vaso-dilatator action, which being thus unmasked becomes evident. In chloral poisoning it is usual for stimulation of afferent nerves to give at once a fall, not a rise, of blood-pressure. It would appear that constricting action is more affected by chloral than dilating action. These results refer to the reflex effects of stimulation of afferent nerves; very similar results are obtained by the direct stimulation of efferent (vasomotor) nerves. The sciatic, for instance, if its peripheral end be stimulated, gives, if fresh, vaso-constriction—but if fatigued, vaso-dilatation. Here again it appears that dilating outlasts constricting action. The effects are such as to remind us of those obtained by prolonged stimulation of the cardiac nerves; when vagus and accelerans are simultaneously excited, the vagus effect wears away comparatively quickly, and the accelerator effect is left unmasked.

*Local vasomotor reflexes.*—As has been stated, the effect of the stimulation of efferent nerves is usually vaso-constriction and rise of general blood-pressure, but sometimes the reverse. The local effects are precisely opposite; usually the circulation



becomes more active in a part of which an afferent nerve is excited, the vessels of the part dilate, while the rise of the general blood-pressure shows that other vascular districts of the body at the same time contract; both factors thus concur in promoting the more copious blood-supply of the district, an afferent nerve of which is stimulated; the effect is produced by experimental stimulation of the central end of a divided nerve containing afferent fibres, or by irritation of the cutaneous periphery, the nerves of which are intact, or by excited action of a part. Irritation of the central end of the great auricular nerve (in the rabbit) and consequent dilatation of the vessels of the ear, irritation of the central end of the tibial nerve and consequent dilatation of the external saphena vein, are the two best-known instances of reflex vaso-dilatation by experimental stimuli; the effect of a blister is a familiar instance of reflex vaso-dilatation by cutaneous stimulation; the more active circulation in contracting than in resting muscle, in secreting than in resting gland, are physiological instances of reflex vaso-dilatation in consequence of increased activity of a part, to which may be added as a pathological instance the increased activity of the circulation in an 'inflamed' part. In all these cases peripheral activity causes centripetal stimulation, the reflex effects of which are vaso-dilatation and increased blood-supply.

It should finally be said that the contractility of blood-vessels—arteries as well as veins—may manifest itself independently of influences through nerves from the central nervous system. As regards veins, the most striking instance is that afforded by the bat's wing (Wharton Jones), in which rhythmic constrictions of the veins may be observed, occurring once or twice per minute, and owing to the presence of valves—promoting the onward course of the blood. As regards arteries, the fact that they can gradually recover 'tone' subsequently to nerve-section shows that tone is not *entirely* of central origin. An extract of the suprarenal gland—of the medulla, not of the cortex—injected into the vascular system causes a well-marked increase of arterial blood-pressure; this increase is due to a direct action upon the arterioles, not a might have been supposed to an indirect effect through vasomotor centres and nerves (Schäfer and Oliver).

#### THE LYMPHATIC CIRCULATION

The lymph moves in a circle, but very slowly in comparison with the blood; it exudes into the tissue-spaces through the

walls of the capillaries, and is carried back to the blood by lymphatic capillaries and vessels, converging from all parts to form the thoracic ducts, which discharge their contents into the subclavian veins. The forces by which the current of lymph is kept up are:—(1) the exudation pressure under which lymph is discharged, and (2) the accidental compression of lymphatic spaces and vessels by muscular movements, and by the arterial pulse throughout the body; the lymphatic vessels are abundantly beset with valves which permit fluid to pass towards the heart, but prevent it from being driven backwards towards the tissues; moreover, the walls of the larger vessels are contractile; a slight favouring action is attributed to the current of blood in the subclavian veins, and the aspiratory action of inspiration must act in the same sense; in the lacteals, in which valves are abundant, the flow of chyle is promoted by the pumping action of the villi and by any movement of the intestine.

With regard to the origin of lymph and the mechanism of its discharge, it is obvious that the blood is the ultimate source of all lymph; from the researches of the Leipzig school, under the guidance of Ludwig, we know that the two factors with which the lymph-flow rises and falls are—(1) blood-pressure, (2) muscular movements. But according to Heidenhain's researches, in addition to these mechanical factors, another variable force must be reckoned with, *i.e.* secretory activity—(1) of the epithelium forming the capillary walls, and (2) of the tissue-cells themselves. Heidenhain has shown further that the lymph secreted directly from the capillaries or indirectly from the tissues can be distinguished and separately influenced by the injection of various substances; *e.g.* extract of crab's muscle favours the secretion of 'blood-lymph,' while urea, or salt, or sugar, favours the discharge of 'tissue-lymph.' One of his most conclusive proofs of lymph *secretion* is afforded by the fact that the flow of lymph in the thoracic duct will continue for a considerable period in spite of occlusion of the aorta and fall of blood-pressure, and that it is increased, in spite of reduced blood-pressure, after occlusion of the vena cava, or of the vena portæ.

In some of the lower animals special contractile organs, the so-called lymph-hearts, by their rhythmic action, forward the movement of the lymph. The frog possesses two such pairs of hearts, an interior pair beneath the scapula, and a posterior pair in the ileo-coccygeal space; their functional relations are



in many respects analogous with those of the blood-heart ; their nervous supply is derived from the second and from the tenth pairs of spinal nerves, and from the sympathetic ; the muscle which enters into their composition is similar to that of the heart ; and although the experimental effects of nerve-section and nerve-stimulation are not so precise as might be desired, we have evidence that the action of the lymph-hearts, while locally independent of the spinal cord, is subject to reflex inhibition like that of the heart itself. The rate of beat, usually ranging from 60 to 80 per minute, is, however, far more variable, and each lymph-heart has its own rhythm independent of that of the others.

Observations on the pressure and flow of fluid in the larger lymph-vessels indicate a far more sluggish circulation than in the case of the blood ; in the thoracic duct the pressure is between 1 and 2 cm. of water, and the current flows at a rate below  $\frac{1}{2}$  cm. per second, and the total amount of fluid traversing the duct per diem is less than 5 litres. In its passage through the lymph-glands the fluid receives newly-formed leucocytes and becomes more highly charged with proteid. The glands just alluded to are essentially masses of lymphoid tissue, bordering upon what is termed the lymph-channel, through which the fluid percolates from afferent to efferent lymphatics of the glands. This lymphoid tissue is the source of leucocytes, and plays the part of a physiological filter in relation to lymphatic absorption ; this is illustrated by the effects of poisoned wounds, such as are frequently received in dissecting or *post-mortem* work ; inflammation set up by septic matter extends along the lymphatics to the lymphatic glands, where it usually remains localised.

The total amount of lymph in the body is estimated at 25 to 30 per cent. of the body-weight, *i.e.* no less than three or four times the amount of blood, but the data upon which the estimate is formed are very imperfect, and the estimate is quoted merely to signify that the amount is very large.

An excessive exudation of lymph not carried back by the lymphatics to the venous system is known as œdema or dropsy ; the accumulation may be general or local, it may occur in the connective tissue or in serous cavities, where it forms the lymphatic effusions already alluded to—hydrocele, pleuritic, and pericardiac fluids.

## CHAPTER IV

## RESPIRATION

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## THE LARYNX

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**Internal and external respiration.**—In common language, the term respiration is used indifferently for the visible act of breathing, and for the invisible gaseous exchanges taking place between the air and a living body. We have at the outset to distinguish in the complete process of respiration two stages: *external*



respiration, or the mechanism of aëration; and *internal* respiration, or the actual exchange of gases that takes place in the tissues of the body. The essential function, which is the property of every living cell, vegetable and animal, of low as of high degree, is internal respiration; external respiration is accessory to it, inasmuch as it is the preliminary mechanism, varying in various cases, by which that essential function is made possible. Thus in the lowest organisms we have the whole process effected by direct diffusion without the intermediation of respiratory or circulatory organs: among higher animals (vertebrata) we have a series of preliminary phenomena effected through a respiratory organ—gills, or skin, or lungs—and through a circulating medium—the blood: by these means oxygen is carried to every living cell of the body, and carbon dioxide is carried away. Strictly speaking, the process is divisible into four stages: (1) the mechanism of breathing, viz. the respiratory movements; (2) pulmonary or external respiration, viz. the exchange of gases taking place between the air and the pulmonary blood; (3) systemic or internal respiration, viz. the exchange of gases taking place between arterial blood and lymph or tissue; (4) the chemical changes which take place in living aërated tissue, *i.e.* the tissue-nutrition of which oxygenation forms part. That these are the links into which the respiration of a mammalian animal is naturally divisible will be clearly brought out by a brief historical sketch of the stages through which our knowledge of the subject has progressed.

I. Black, in 1757, showed that the end product of respiration is identical with that of combustion, viz. carbon dioxide; Priestley, in 1775, discovered oxygen, compared respiration with combustion, and contrasted the respiration of plants with that of animals. Lavoisier definitely established the fundamental facts, viz. that respiration, like combustion, consists in consumption of oxygen with production of carbon dioxide, and evolution of heat.

II. Lavoisier's followers believed that the production of carbon dioxide was the immediate consequence of the consumption of oxygen, and taught that respiration was a direct oxidation or union of oxygen with carbon taking place in the lungs.

III. That this is not the case was proved by the extraction of the blood-gases by Magnus in 1837. The fact that oxygen and carbon dioxide exist as such in the blood proves that the lung at any rate is not the seat of oxidation.

IV. Nor is it the blood (as was for a time supposed), for carbon

dioxide was extracted by Pflüger and his pupils from the lymph and from muscular tissue.

V. Thus far the process is traced back to its source in the living tissues; but now, finally, it has been shown in the laboratories of Pflüger, of Ludwig, and of Hermann, that the process itself is no direct union of oxygen with carbon, but that oxygen is first integrated in some complex body, by the subsequent disintegration of which carbon dioxide is produced. That carbon dioxide is not the *immediate* consequence of oxygen supply is demonstrated by the fact that living tissue, or even an entire animal, such as a frog, goes on exhaling carbon dioxide in an atmosphere free of oxygen.

Thus the links in the chain of respiration, as we now know them, are as follows:—*Oxygen, introduced into the lung by muscular movement, diffuses into the pulmonary blood and is conveyed to the systemic capillaries, whence it diffuses into the lymph and tissues; here it enters and forms part of some complex compound, which subsequently yields carbon dioxide as a disintegration product; carbon dioxide diffuses from the lymph to the blood, and is therein carried to the lung, whence it diffuses into the air.*

**Physiological anatomy of the lungs and thorax.**—The thorax and lungs together constitute a bellows, by the alternate enlargement and shrinkage of which air is drawn in and driven out through the nostrils, windpipe, and air-passages. The lungs are practically a many-chambered bag contained in the thorax, not attached to its walls, but in close apposition to them; the outer surface of the lungs and the inner surface of the thorax are covered by a serous membrane, the *pleura*; between these two layers is the pleural cavity, which in health is practically non-existent, inasmuch as the two layers are in close apposition, moistened by a barely appreciable amount of lymph. Lungs and thorax move conjointly, expansion and contraction of the thorax entailing expansion and contraction of the lungs; when the thorax and lungs expand, air enters the lungs by atmospheric pressure; when the thorax contracts, air is driven out into the atmosphere.

The thorax is a closed cavity; if it should be broken into by accident or by operation, the lung does not remain in apposition with the chest-wall, but shrinks away from it and collapses, while the chest-wall, relieved of the elastic traction normally exercised by the lung, undergoes a slight expansion. If the thorax should be perforated, the lung in collapsing separates the two layers of the *pleura*, and the pleural cavity is occupied by air



(pneumothorax). If the membrane should become inflamed, lymph and leucocytes are effused into the pleural cavity (pleuritic effusion), and as one of the after-effects of inflammation it may happen that 'adhesions' are formed between the two layers of the pleura, leading to a more or less complete obliteration of the pleural cavity. Collapse of both lungs causes immediate death, collapse of only one lung is dangerous but not necessarily fatal. During foetal life the lung contains no air and exercises no elasticity (atelectasis); after birth, with the establishment of pulmonary respiration, the alveoli are opened out, and by the subsequent growth of the thorax the lungs undergo a further passive expansion.

The lung ultimately consists of an aggregation of minute chambers—the *alveoli* or *air-cells*—the walls of which are covered with a close network of capillaries. This subdivision of the air-sac affords a very extensive surface exposed to the air which penetrates to the alveoli; and the capillary network affords a similar spreading-out of the blood into what amounts to a thin sheet of very great area. It has been estimated that the alveolar surface amounts to no less than 200 square meters, and that the total capillary surface is equal to 150 square meters; the thickness of the capillary sheet of blood may be reckoned as being somewhat greater than that of a single blood-corpuscle, say  $10\mu$ . From these data it may be realised how favourable are the conditions to the prompt aëration of the blood by the lung. At any moment it contains in its capillaries nearly 1,500 c.c. of blood spread out in a thin sheet 150 square meters in area, and only  $\frac{1}{100}$  millimeter in thickness. This extensive film of blood is separated from the alveolar air by the endothelium, which lines the pulmonary capillaries, and by that which lines the inner surface of the alveoli. Through this septum the exchange of gases between air and blood is effectually and rapidly carried out.

The passages through which air enters the lungs are: the nostrils (properly), or mouth (improperly), the trachea, and the bronchi—large and small. The smallest bronchi (1 mm. in diameter) open out into the clusters of air-cells called the infundibula. As regards the structure of these parts, the points of most physiological importance are:—(1) that the epithelium lining the air-passages is of the columnar, ciliated kind, the sharp stroke of the cilia being in the outward direction; (2) that the plain muscle forming part of the tissues of which the air-tubes are composed

is relatively most developed in the smallest bronchi; and (3) that the lumina of all the larger air-tubes are kept open by more or less complete rings and plates of cartilage.

Each act of respiration is composed of two phases—**inspiration** by expansion of the thorax and lungs, **expiration** by their shrinkage. Normally, inspiration is effected by muscular action, while expiration is literally a ‘shrinkage’ to the original volume by virtue of the elasticity of the lungs. It is only when respiration is forced or laboured that expiration is assisted by the action of thoracic muscles. The muscles acting in *normal inspiration* are the diaphragm and external intercostals with the scaleni, the inter-cartilaginous portion of the internal intercostals, the levatores costarum, and the quadratus lumborum. In *laboured inspiration* other muscles come into action—the serrati (magnus, superior, and inferior), the sterno-mastoid, and, indeed, any muscle extending between the thorax and upper extremities, *e.g.* pectorales (major and minor), latissimus dorsi. These last-named muscles, which ordinarily act from the thorax as a fixed point of origin, now act upon the thorax, fixed objects being grasped with the hands so that the humerus and scapula afford points of origin instead of points of insertion. At the same time the vertebral column is fixed and extended by the dorsal muscles. In *normal expiration*, as already stated, the chief factor is the elastic recoil of the lung to its position of rest, with perhaps some slight assistance by the interosseous portions of the internal intercostals and by the triangularis sterni. In *laboured expiration* the abdominal muscles assist by fixing and compressing the abdomen, forcing the diaphragm upwards, and constricting the lower part of the thorax.

The most important of the muscles above enumerated are those which always act in respiration, *i.e.* those of normal inspiration, viz. the diaphragm and the external intercostals (in conjunction with the scaleni). The diaphragm is a dome-shaped sheet of muscle forming the partition between thorax and abdomen. The external intercostals form two lateral sheets of muscle composed of a series of slips extending from rib to rib. The scaleni fix the first two ribs, thus affording an essential condition of effective action as regards elevation of the ribs by the series of external intercostals; each of these muscles acts from the rib above as its relatively fixed point upon the rib below as a relatively movable lever which it elevates; the quadratus lumborum, acting from the pelvis upon the last rib, affords a fixed point to



the contracting diaphragm. By these means the thorax is enlarged in all its diameters—(1) vertically by the descent of the diaphragm, (2) laterally by the elevation of the ribs, and (3) in an antero-posterior diameter by the elevation of the ribs and of the sternum. And according as one or other of these muscular agencies takes chief part in such inspiratory movement, two types of respiration are distinguished—the costal or thoracic, and the diaphragmatic or abdominal. In the thoracic type, which is characteristic of women, the thoracic muscles play the greater part; in the abdominal type, which is characteristic of men, the diaphragm is comparatively more effectual. But it is a mistake to suppose

that these are fundamental sexual differences: they are probably due to differences of dress.

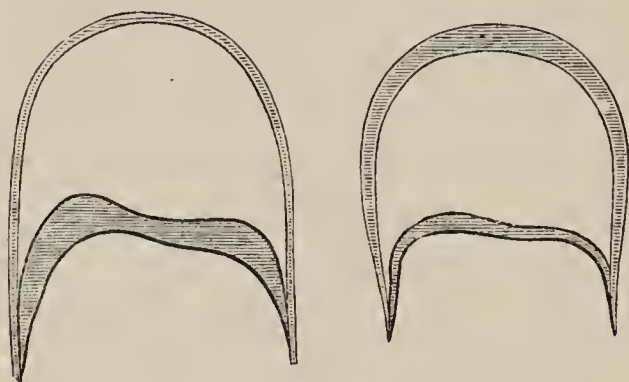


FIG. 57.

To illustrate the 'diaphragmatic' and 'costal' types in the male and in the female. The shaded spaces are intended to indicate the range of movement (much exaggerated) of the diaphragm and of the chest in the two cases.

In ordinary easy breathing a moderate amount of air is taken into the chest with each inspiration, and given out with the succeeding expiration. This air is called the *tidal* air, and amounts to about  $\frac{1}{2}$  litre, or 500 c.c., or 30 cubic inches. Beyond this ordinary or tidal amount it is

possible by an extraordinary effort of inspiration to introduce into the chest a further 1,500 to 2,000 c.c. of air, which amount is termed *complemental* air, or by an extraordinary effort of expiration to expel 1,500 c.c. (in excess of the normal tide), which is termed the *supplemental* air. After the most complete possible expiration, there is left in the lungs a quantity of air amounting to another 1,500 c.c., which no effort can expel; this is the *residual* air. In relation to this matter, two other terms require to be defined—the *vital capacity*, amounting to between 3,000 and 4,000 c.c., is used to denote the amount of air which can be given out by the deepest possible expiration after the deepest possible inspiration; the *capacity of equilibrium* or *stationary air*, amounting to about 3,000 c.c., is the cubic capacity of the chest after normal expiration. These terms will be best brought to mind by referring to the diagram (fig. 58).

We have seen that at each respiration the lungs are not entirely emptied of and refilled with air, but that the greater propor-

tion of their contents remains stationary, while a small proportion only, nearest to the outlet (nose and mouth), is actually exchanged. It is by the rapid diffusion of gases taking place between the stationary and the fresh tidal air that the former discharges carbon dioxide and is replenished with oxygen. We have now to consider in detail the manner in which the pulmonary blood and air influence each other. The effect of the blood upon the air is known by comparing expired with atmospheric air; the effect of air upon blood by comparing pulmonary venous with pulmonary arterial blood.

**Expired air as compared with atmospheric air** contains about 5 per cent. less oxygen and 4 per cent. more  $\text{CO}_2$ . It is warmer,

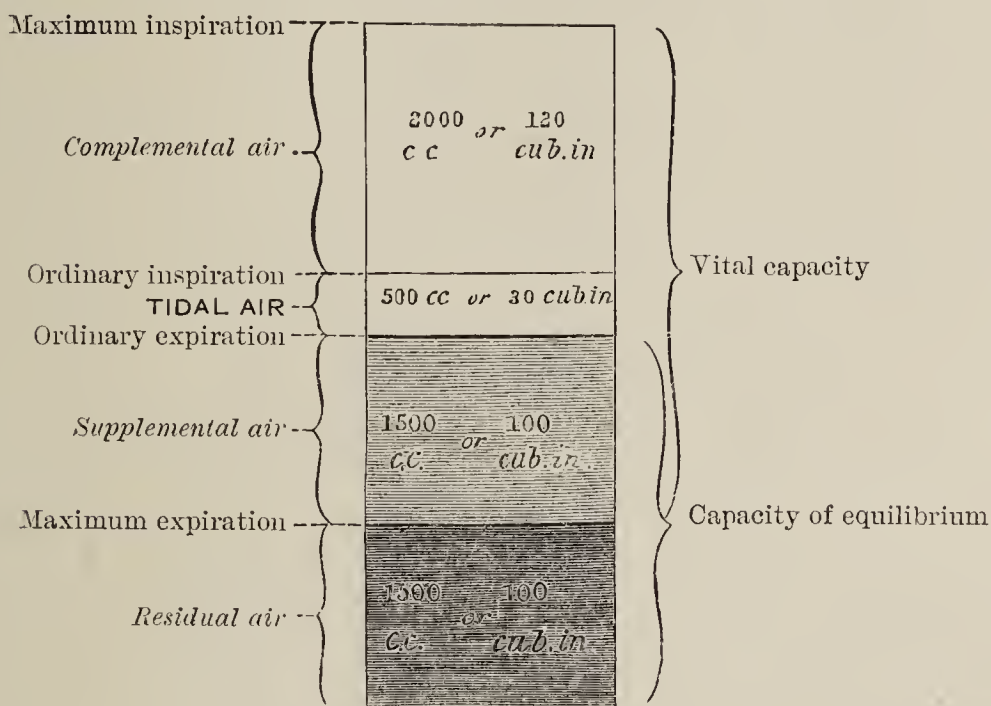


FIG. 58.

Amounts of air contained by the lungs in various phases of ordinary and of forced respiration.

saturated with moisture, fouled by organic emanations, and slightly diminished in volume.

The differences between pulmonary venous and pulmonary arterial blood must obviously correspond with these—theoretically at least—seeing that any gained or lost matter in expired air must, regarded from the other side, be lost or gained matter from the pulmonary venous blood. Practically, however, we can only say that the difference between blood in the pulmonary artery and in the pulmonary veins is that the former is blacker, contains more carbon dioxide and less oxygen than the latter; in a word, that the former is ‘venous’ in character, the latter ‘arterial.’ We cannot state, as directly demonstrated, the minute differences which must exist, either in temperature or in amount of

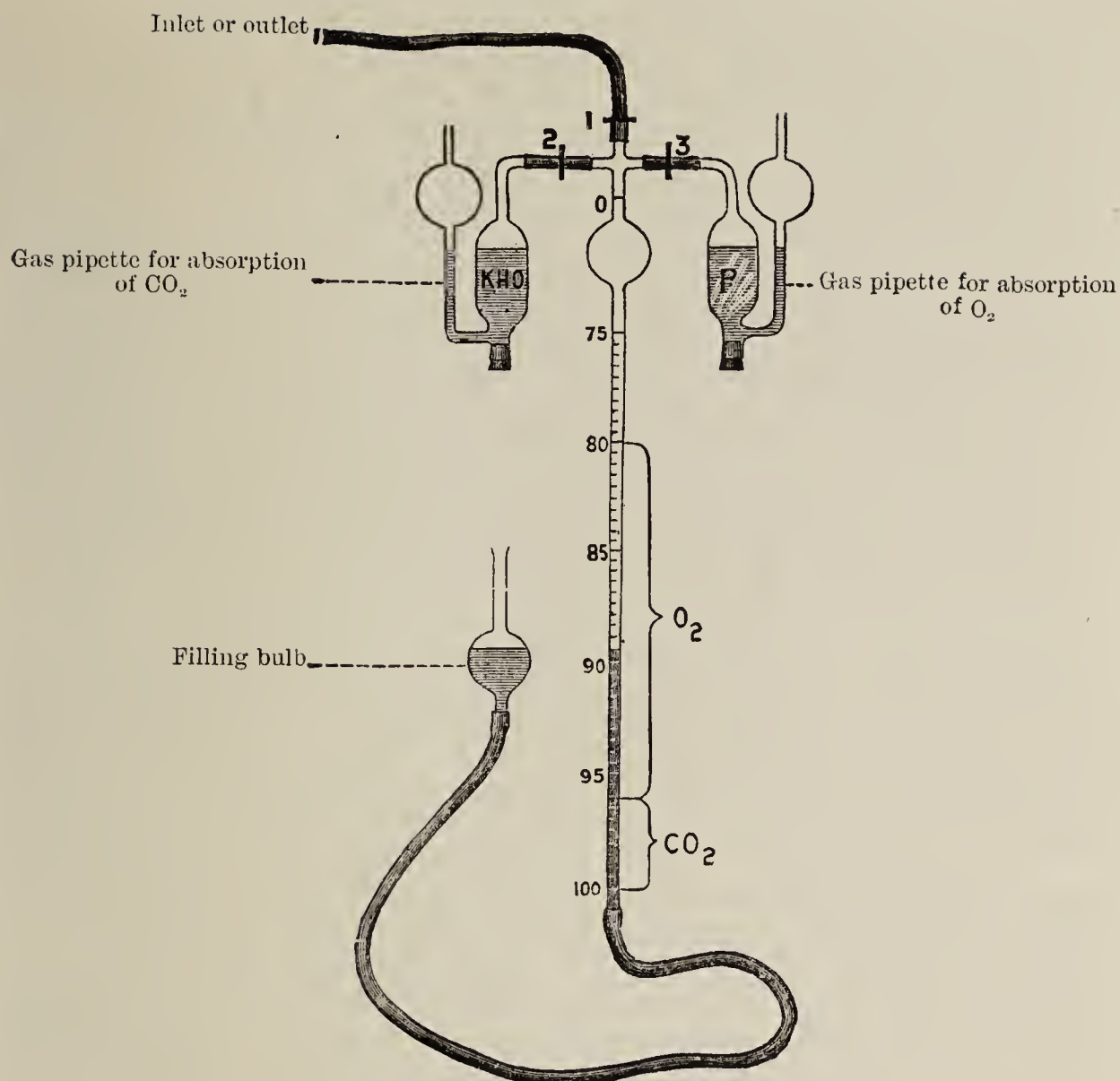


water, or in amount of organic matter. To return to the differences between expired and atmospheric air, it is to be observed that the minus quantity of oxygen is greater than the plus quantity of carbon dioxide; with this inequality corresponds the fact that the volume of expired air is slightly less than that of the previously inspired air. The fraction denoting the ratio  $\frac{\text{Vol CO}_2 \text{ exhaled}}{\text{Vol O}_2 \text{ absorbed}}$  is spoken of as the *respiratory quotient*; normally this fraction is about  $\frac{4}{5}$  or  $\cdot 8$ .

If for every five volumes of oxygen absorbed only four volumes of CO<sub>2</sub> are exhaled, one volume of oxygen remains unaccounted for; it is probable that this excess of oxygen which does not appear in union with carbon does so in union with hydrogen as water.

A second point, which is of great practical importance, relates to *organic matter* exhaled from the lungs. It is the chief factor in the fouling of air, but, though its effect is thus so pronounced, no direct estimate or measurement of its amount can be made. All we may say concerning its nature is, that it is probably proteid in character, condensed respiratory moisture giving a faint xanthoproteic reaction. It may, however, be indirectly estimated by measurement of exhaled CO<sub>2</sub>, the amount of which indicates to what degree the air is fouled by the more deleterious but impalpable organic emanations of the breath. A rough but excellent guide to the amount of respiratory impurity is the sense of smell. The air of a room in which respiratory CO<sub>2</sub> does not exceed 2 per 10,000 is 'fresh'; when the respiratory CO<sub>2</sub> is between 2 and 4 per 10,000 the room begins to feel close, between 4 and 6 per 10,000 it is decidedly 'close,' between 8 and 10 the air is 'foul,' and beyond this limit intolerable for any length of time. 2 per 10,000 or 1 per 5,000 is thus the limit of expired CO<sub>2</sub> admissible in perfect ventilation, and should never be exceeded. But although this comparatively high standard of purity is desirable in the air of permanently occupied chambers, we frequently, and without discomfort, remain for short periods in air with a much higher percentage of respiratory CO<sub>2</sub>—*e.g.* in theatres or in class-rooms. CO<sub>2</sub> is not of itself specially injurious—an ordinary gas-light gives off nearly as much CO<sub>2</sub> as ten people, but the air—although spoiled to some extent—is not nearly as much spoiled by one gas-burner as by ten people; thus, in gauging chamber-air it is mainly the *respiratory* CO<sub>2</sub> which should be estimated.

From the data given above we may at once calculate the

FIG. 59.—ESTIMATION OF  $O_2$  AND OF  $CO_2$  IN EXPIRED AIR.

A 100 c.c. measuring tube graduated in tenths of a c.c. between 75 and 100. A filling bulb. Two gas pipettes. The measuring tube communicates with the inlet and with the gas pipettes by three tubes guarded by clips 1, 2, 3. It is first charged with acidulated water up to the zero mark by raising the filling bulb, tap 1 being open; it is then filled with 100 c.c. of expired air, the filling bulb being lowered until the fluid in the burette has fallen to the 100 mark. Tap 1 is now closed, the measuring tube containing 100 c.c. of expired air with unknown quantities of  $CO_2$  and of  $O_2$ . The amount of  $CO_2$  is ascertained as follows: Tap 2 being opened, the air is expelled into a gas pipette containing KHO by raising the filling bulb until the fluid has risen to the zero mark of the measuring tube. Tap 2 is now closed, and the air left in the gas pipette for about a minute, during which the  $CO_2$  present is entirely absorbed. The air is then drawn back into the measuring tube by lowering the filling bulb while tap 2 is open. The volume of air (minus the  $CO_2$ , which is absorbed) is read, the filling bulb being adjusted so that its contents are at the same level as the fluid in the burette. The amount of  $O_2$  is next ascertained in a precisely similar manner by sending the air into a second gas pipette containing sticks of phosphorus in water, and measuring the loss of volume (due to absorption of  $O_2$ ) in the air when drawn back into the tube. A gas pipette works thus: fluid in its lower half is displaced into its upper half when air is driven in from the measuring tube, and returns to its original place when air is drawn back. If desired, the apparatus can be connected with a vessel in which a frog, or mouse, or excised muscle has been placed, and the consequent alterations of the gases  $O_2$  and  $CO_2$  measured in a similar manner.

The analysis of a single sample of expired air is not a reliable indication of respiratory activity. It is necessary to know the total amount of air expired in a given time, and to analyse a mixed sample of that total amount corrected to standard temperature and pressure. These conditions are fulfilled by Zuntz'



complete apparatus. A gasometer records the total amount of air expired, and at the same time unwinds a filling apparatus so as to collect in a measuring tube an average sample of the total air expired. If, for instance, during 5 minutes, 40 litres of air were expired, and if an average sample of 100 c.c. collected during that period contained 4 c.c. more  $\text{CO}_2$  and 4.5 c.c. less oxygen than atmospheric air, the average per minute would be 320 c.c.  $\text{CO}_2$  discharged, 360 c.c.  $\text{O}_2$  absorbed, and the respiratory quotient =  $\frac{8}{9}$  or .88.

To examine the respiratory exchange of animals it is necessary to connect the gasometer with a tube tied into the trachea, or with an air-tight 'respiration chamber' in which the animal or man is confined; in this case, air is drawn through the chamber and gasometer by an aspirator. The older apparatus of Pettenkofer and Voit, as well as that of Regnault and Reiset, was constructed upon this last principle, including (1) a respiration chamber, (2) a gasometer, (3) an aspirator, and (4) a series of baryta tubes in which excreted  $\text{CO}_2$  was absorbed and weighed, the oxygen item being undetermined. For experiments extending over long periods, the total amount of  $\text{CO}_2$  excreted by a large animal is inconveniently large; the difficulty is met by taking only a measured fraction of the total expired air through the absorption tubes to be weighed.

*amount of fresh air* which should circulate through a properly ventilated room, remembering that every 1 c.c. of expired  $\text{CO}_2$  is to be diluted by at least 5,000 c.c. of fresh air. An adult expiring 15 times per minute 500 c.c. of air containing 20 c.c. of  $\text{CO}_2$  exhales in one hour 18 litres of  $\text{CO}_2$ , which, to be diluted 5,000 times, requires 90,000 litres of air, *i.e.* the fresh air required during the hour is 90,000 litres, or 3,000 cubic feet. This is the normal amount of fresh air per head per hour which should be provided for in the ventilation of dwellings intended to be 'healthy.'

**Frequency and depth of respiratory movements.**—We have taken as an average frequency of respiratory movements 15 per minute. The frequency is by no means invariable; anything between 12 and 20 is normal, and the number is subject to many modifying

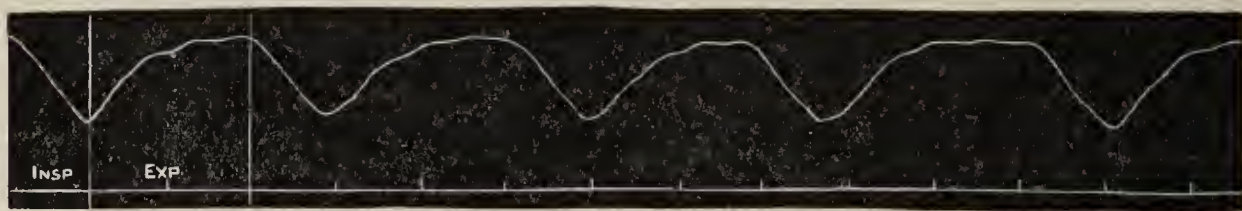


FIG. 60.

Tracing of normal respiration of man. The frequency of the act was at the time 20 per minute; each respiration occupied 3 sec., of which inspiration occupied 1 sec., expiration (inclusive of the expiratory pause) 2 sec. The smaller undulations (about 4 to each respiration) are caused by the heart-beats.

conditions—as age, sex, exercise, health—and also to various emotional and mental states. The frequency of respiration usually bears a definite ratio to pulse-frequency, *viz.* 1 respiration to every 4 to 5 heart-beats. It is greater in the female or child than in the male or adult. It is increased by exercise, and diminished during quiescence, but variations of depth are more

important than variations of frequency. During illness, especially in the febrile state, it is increased. As regards the effects of emotional and mental states, we shall more conveniently consider them under the heading 'Influence of the Nervous System'; it need here only be observed that the frequency of respirations should always be noted while the patient is unaware of the proceeding, and not under the influence of any exciting event, because any disturbing cause—the mere fact of turning attention to the act of breathing—may suffice to alter its character and frequency.

An accelerated rate of respiration is accompanied with an increased elimination of  $\text{CO}_2$ , but this increased elimination is by no means proportional to the greater frequency of respiration; this is accounted for by the fact that each act of respiration is deeper in a slow series than in a rapid series, so that a man breathing 30 times per minute does not expire anything like twice as much air, nor twice as much carbon dioxide, as a man breathing 15 times per minute.

The percentage of  $\text{CO}_2$  is less in the expired air of shallow than in that of deep respiration; it is less in the air expelled at the beginning than in that of the end. By holding the breath the percentage may be raised from 4 up to, but not beyond, 8 per cent. It is of importance to distinguish the separate effect of increased frequency and of increased depth upon the total discharge of  $\text{CO}_2$ . We should have the same amount of air expired per minute (8 litres) in these two cases:  $\text{depth} = 500 \text{ c.c.} \times \text{frequency} = 16$ , or  $\text{depth} = 400 \text{ c.c.} \times \text{frequency} = 20$ , but in the first case the  $\text{CO}_2$  exhaled would be greater than in the second. From which we may realise that to carry off the product of a more active internal respiration, *deeper* breathing is more effectual than *more frequent* breathing; and as a matter of fact, variations of activity affect the depth more than they do the frequency. Muscular exertion may increase the frequency, but it more prominently increases the depth of respiration, and there can be little doubt that *plus*  $\text{CO}_2$  in the blood is the excitant of the deeper breathing by which *plus*  $\text{CO}_2$  is discharged; one may experience for oneself that an addition of  $\text{CO}_2$  to air inspired from the gas-holder causes a deeper respiration without discomfort, or even any knowledge that a change has occurred. In rapid, shallow breathing the discharge of  $\text{CO}_2$  is comparatively low, and it is the amount of evaporation that is principally augmented. This type of respiration occurs when the temperature of the blood is raised, or tends to



rise, and the increased evaporation has a cooling effect. Dogs breathe thus when hot; frequent shallow respiration is characteristic of fever; and the so-called '*heat-dyspnœa*,' that can be experimentally produced by warming the carotid blood, is of a similar type—frequent and shallow.

*Dyspnœa. Asphyxia. Apnœa.*—Dyspnœa signifies difficult breathing, and is the consequence of any impediment to the free ingress and egress of air from the lungs. If aëration be com-

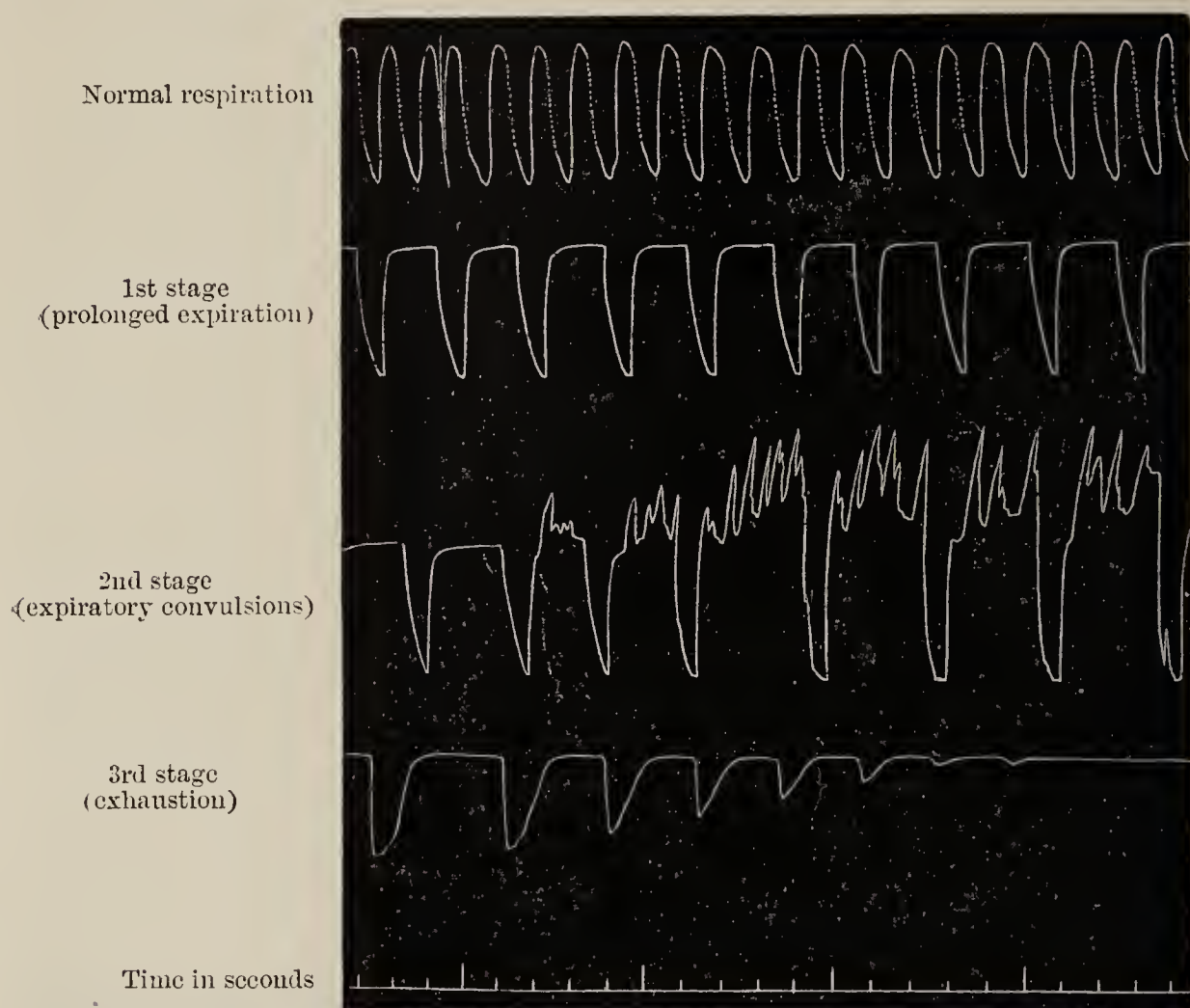


FIG. 61.—ASPHYXIA TRACING. (Rabbit.)

(The line falls with inspiration, rises with expiration.)

pletely interfered with, as by drowning or by obstruction of the trachea, death by asphyxia is the result. The term asphyxia, which literally signifies pulselessness, is generally taken to cover the entire series of events from the moment when aëration is interfered with, to the cessation of all respiratory and cardiac movements. This series of events, as observed upon an animal killed by sudden and complete closure of the trachea, is divisible into three stages: (1) that of increasing dyspnœa, culminating in (2) the convulsive stage, which gives way to (3) the period of exhaustion. During the first stage, which lasts about a minute, the respiratory movements become stronger and longer, therefore

less frequent, the prolongation being chiefly expiratory; the heart beats with increased force and frequency, the blood-pressure rises, the pink or red colour of the tongue and lips darkens and gives place to bluish purple. *During the second stage*, which lasts about a minute, the respiratory movements become violent and convulsive, the convulsions being chiefly expiratory; the heart beats forcibly but less frequently, the blood-pressure keeps high, the dusky colour of the mucous membranes is increased. That the raised blood-pressure is caused by vaso-constriction rather than by increased heart's force is proved by the fact that the blood-current is at the same time much slower than normal. *During the third stage*, which lasts two or three minutes, movements ebb, the pupils dilate, convulsions cease, respiration becomes less frequent and more shallow, finally ceasing in expiration; the heart beats feebly and unfrequently, or feebly and rapidly, and finally stops, the blood-pressure falls and nearly reaches zero, its decline being frequently marked by Traube-Hering undulations; the dusky mucous membranes become pale and anæmic. *Post mortem* the right side of the heart, the large veins and the lungs are gorged with thick blood, the left side of the heart is empty and contracted.

Recovery from asphyxia is possible in apparently almost hopeless cases; artificial respiration, *i.e.* rhythmical inflation of the chest by means of bellows (on animals), or passive movements of the arms and chest in imitation of inspiration and expiration (on man), will often cause the heart to resume action, and blood-pressure to be restored; relief of the engorged right auricle by venesection will sometimes set going again a heart that seems to have finally stopped. The practical lesson which these facts enforce is that artificial respiration should be steadily maintained on the bodies of the apparently drowned, and that a jugular vein, especially if visibly distended, may be and should be opened. Artificial respiration is also in common use in the laboratory to keep animals alive when from any cause their spontaneous movements of respiration can no longer be effected. Poisoning by carbon monoxide is practically death by asphyxia, the hæmoglobin is fixed as carboxyhæmoglobin and rendered incapable of acting as an oxygen carrier.

*Apnœa* is the opposite of dyspnœa: it is the temporary non-breathing state which is induced by repeated inspiration or by inflation of the lungs. An animal rendered apnœic by artificial respiration remains passive for several seconds without any effort



of respiration, and then begins to breathe by very shallow inspirations, which gradually deepen. This passive state may be brought about on oneself by repeated deep inspirations, after which it will be found that the breathing may be suspended for a longer period and with greater comfort than in the normal state. The apnœic state has received more than one explanation ; it has been referred to a more completely saturated state of arterial blood as regards oxygen, to a provision in the lung itself of a larger and purer air-supply, and to an inhibitory effect produced by the mechanical distension of the lungs. Probably all these factors concur to the result, for these reasons :—apnœa in animals is *best* produced by inflation with oxygen, but *can be* produced by inflation with hydrogen or nitrogen ; apnœa is *best* produced with the vagi nerves intact, but it *can be* produced after section of both these nerves. Apnœic pauses on man are longer and shorter according as the amount of oxygen inhaled is greater or smaller, but in no constant ratio with that amount ; the non-correspondence of the two quantities is attributable to the fact that nervous influences are also in play in the maintenance of apnœic pauses : these nervous influences are probably (1) an inhibitory effect transmitted along the vagus to the spinal bulb from the distended lung, and (2) in a minor degree a depression of the excitability of the ‘respiratory centre’ by blood fully charged with oxygen.

The immediate cause of dyspnœa is principally a deficiency of oxygen, but it may also be—and under normal circumstances probably is—provoked by an accumulation of  $\text{CO}_2$ . If, experimenting upon oneself, one breathes in and out of a gas-holder containing pure hydrogen instead of air, dyspnœa is quickly developed ; this is due to want of oxygen. If one breathes pure oxygen, dyspnœa is more slowly developed ; this dyspnœa is due to excess of  $\text{CO}_2$ , for it is produced long before the oxygen percentage in the gas-holder is reduced to 20 per cent. What is known clinically as ‘cardiac dyspnœa’ is essentially an oxygen dyspnœa, the supply of oxygenated blood is barely adequate to the needs of the quiescent body, and, owing to the imperfect action of the heart, increased demand, by even slight exertion, fails to be met by increased supply, the respiratory centre becomes excited and dyspnœa is witnessed, which is exactly similar to the ‘breathlessness’ and violent breathing that a normal person manifests after more vigorous exertion. ‘Hæmorrhagic’ dyspnœa is likewise due to a lack of oxygen caused by loss of blood.

*Cheyne-Stokes breathing* is characterised by a waxing and

waning of the amplitude of the respiratory movements. In a typical and well-marked case, the movements alternately decline to complete cessation, and return to an amplitude much above the normal. No definite or conclusive cause can be assigned to this peculiarity of rhythm; it is not, as was once believed to be the case, characteristic of fatty degeneration of the heart, but makes its appearance in a variety of diseases, or in the absence of any disease at all; during normal sleep, particularly in children, a waxing and waning respiratory rhythm is of common occurrence. All we can say in explanation is to point to the fact that the Cheyne-Stokes rhythm is to the respiratory system what the Traube-Hering rhythm is to the vasomotor system; both rhythms are originated by bulbar centres and are of about

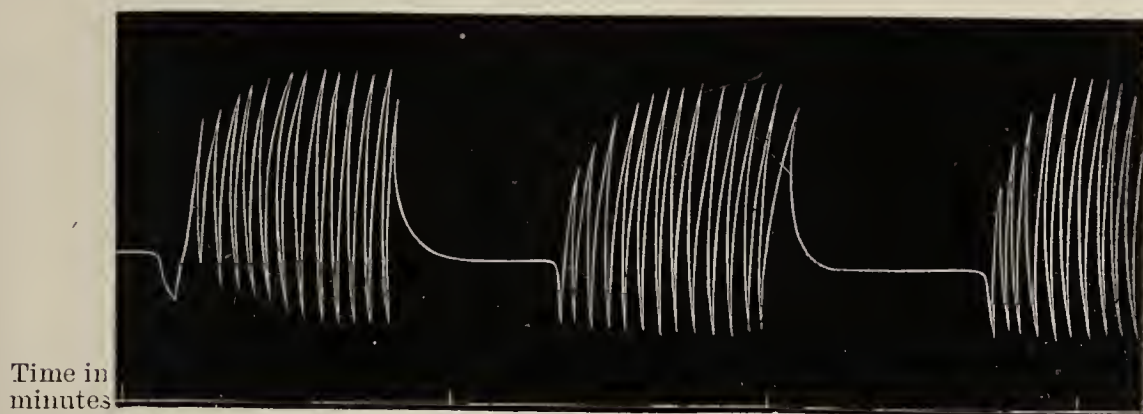


FIG. 62.—CHEYNE-STOKES RESPIRATION.

the same frequency, viz. 1 to 3 per minute; indeed, the association is sometimes most definite and exact; on the rabbit, for instance, after hæmorrhage, phases of increasing and diminishing amplitude of respiration coincide with rise and fall of arterial blood-pressure; they are instances among many others of the common tendency towards 'pulsatile or rhythmic activity' manifested by all living matter. The light is apt to flicker, and especially so when it is going out.

**Variations of respiratory activity under different conditions.**—We have learned that the depth and frequency of respiratory movements, and consequently the amount of  $\text{CO}_2$  excreted, vary under different conditions; these variations of the external yield are the consequence of variations of tissue-activity, and therefore afford an index to the variations of the true or internal respiratory activity of the entire body. It should be clearly recognised that the magnitude of the external respiratory exchange is determined by the degree of internal respiratory activity, and that the converse event—a modification of internal by external respiration—is comparatively slight or of accidental occurrence. We shall



take as our point of departure and standard of reference the average respiratory activity of a normal adult. Given—that the average frequency of respirations is 15 per minute, that the average amount of tidal air is 500 c.c., and that expired air contains 5 volumes per cent. less oxygen and 4 volumes per cent. more carbon dioxide—a simple calculation gives the average hourly or daily absorption of oxygen and exhalation of carbon dioxide as follows :

	Volume of air expired	CO <sub>2</sub> exhaled		O <sub>2</sub> absorbed	
		by volume	by weight	by volume	by weight
Per respiration .	500 c.c.	20 c.c.	0.39 grm.	25 c.c.	0.36 grm.
Per minute (15 r.).	7.5 litres	300 "	5.91 "	375 "	5.37 "
Per hour .	450 "	18 litres	35.5 "	22.5 litres	32.2 "
Per day (24 h.) .	10,800 "	432 "	852 "	540 "	773 "

Thus a normal adult (weighing 70 kg.) excretes *per diem* 432 litres CO<sub>2</sub> (weighing 852 grammes, and containing 232 grammes of carbon); he absorbs *per diem* 540 litres O<sub>2</sub> (weighing 773 grammes). Or, otherwise expressed, the average excretion of CO<sub>2</sub> is  $\frac{1}{4}$  litre (weighing  $\frac{1}{2}$  gramme) *per kilo per hour*, or about 4 c.c. per kilo per minute.

It should be particularly observed that these are average numbers, subject to considerable variation. Thus Pettenkofer and Voit's man, weighing 70 to 73 kilos, yielded daily amounts of CO<sub>2</sub> fluctuating between 695 and 1,038 grammes. Moreover, they are subject to considerable fluctuations in varying conditions of health and of activity; thus an excretion of CO<sub>2</sub> between 3 and 5 c.c. per kilo per minute, *i.e.* 180 to 300 c.c. per kilo per hour, may be regarded as normal.

The respiratory activity of different animals under different conditions is usually estimated by measuring the excretion of CO<sub>2</sub>

Animal	Conditions	CO <sub>2</sub> exhaled per kilo per hour
Frog . . . .	Temp. 10° to 15°	45 c.c.
" . . . .	" 30° to 35°	317 "
Hen . . . .	—	787 "
Hen's egg . . . .	during incubation	18 "
Marmot . . . .	hibernating	19 "
" . . . .	awake	667 "
Sheep . . . .	—	350 "
Sheep foetus . . . .	at term weighing 3.5 kg.	30 "
Man . . . .	—	300 "
Rabbit . . . .	—	550 "
Guinea-pig . . . .	—	750 "
Horse . . . .	at rest	200 "
" . . . .	at work	1,200 "

rather than the absorption of  $O_2$ ; for observations in which it is desired to ascertain the respiratory quotient (p. 124) both measurements must be simultaneously carried out.

The subjoined tables contain illustrative numbers from which we may draw certain further conclusions.

The respiratory activity of cold-blooded is less than that of warm-blooded animals. In cold-blooded animals it varies with temperature, being much greater at a high than at a low temperature; in warm-blooded animals the relation is more complex; so long as their body temperature remains constant, their respiratory activity diminishes with a rise and increases with a fall of temperature; but if their body temperature is raised or lowered, then the respiratory activity, as measured by  $CO_2$ , varies in the same sense, being greater at a high than at a low temperature; with the high temperature of fever the excretion of  $CO_2$  is increased, but the respiratory quotient is not diminished. The variations with age, food, and muscular exercise are particularly instructive. During incubation or pregnancy, and immediately after birth, the respiratory activity is low—new-born animals resist asphyxia for many minutes; during infancy and childhood respiratory activity is greater than during adult life; during adult life it is greater than in old age; the average excretion of  $CO_2$  per kilo per hour being—

<i>In utero</i>	.	.	.	.	.	.	.	.	30 c.c.
In childhood	.	.	.	.	.	.	.	.	500 „
In adult life	.	.	.	.	.	.	.	.	300 „
In old age	.	.	.	.	.	.	.	.	250 „

*Muscular exercise* at once raises the excretion of  $CO_2$  above the normal of ordinary life; perfect quietude, on the contrary, lowers it; the excretion of  $CO_2$  by an adult per kilo per hour averages—

During sleep	.	.	.	.	.	.	.	.	200 c.c.
„ quietude	.	.	.	.	.	.	.	.	250 „
„ ordinary exertion	.	.	.	.	.	.	.	.	300 „
„ hard labour	.	.	.	.	.	.	.	.	500 „ or more.

This relation is confirmed by experiments on animals; the  $CO_2$  excreted by a dog or rabbit with the posterior extremities paralysed by division of the spinal cord is below normal; that of the same animal when the posterior extremities are tetanised is above normal. Again, the respiratory excretion of  $CO_2$  by a curarised animal is diminished, that of a strychninised animal is increased. The diminished production of  $CO_2$  from paralysed as



compared with quiescent muscle will be referred to again when we come to consider the respiration of muscle.

*Food* increases the production of  $\text{CO}_2$ , carbohydrates being particularly effective in this direction. The excretion of  $\text{CO}_2$  is greater after than before a meal. Light is favourable to respiratory activity ; it has been shown by experiments on frogs, birds, and mammals, that the excretion of  $\text{CO}_2$  is greater in the light than in darkness, even if the animals have been blinded. The difference is due to an accelerated or retarded tissue-change, as illustrated in the fattening of cattle ; the balance of evidence is to the effect that fattening proceeds better in dimly-lighted than in well-lighted stalls.

Variations in the respiratory quotient are of some theoretical importance, but are difficult to determine with accuracy. From the numbers given below it will be seen that the ratio is nearest to unity with carbohydrate diet, and furthest from it in the hibernating condition, when the animal's chief food is its own fat.

Animal	Condition	Respiratory Quotient $\frac{\text{CO}_2}{\text{O}_2}$
Rabbit . . . .	Fed on carrots	·9 to 1
" . . . .	Fasting	·7
Dog . . . .	Flesh diet	·7
" . . . .	Bread and fat	·9
" . . . .	Fasting	·7
Marmot . . . .	Sleeping	·5
" . . . .	Awake	·8
Man . . . .	Mixed diet	·8 to ·9

It is also increased by muscular exercise, and diminished during quiescence : the respiratory quotient of a rabbit, for instance, is raised by tetanisation, depressed by curarisation. According to Pettenkofer and Voit, it is greater during the day than during the night, daytime being favourable to discharge of  $\text{CO}_2$ , night-time to absorption of  $\text{O}_2$ .

**Respiration of muscle.**—Direct data relating to internal respiration are most readily obtained from muscle, and the following plans have been followed :—1. Examination of the gaseous exhalation of muscle enclosed in air, or in an indifferent gas such as nitrogen, or *in vacuo*. 2. Extraction of the gases of muscle itself. 3. Examination of the gaseous exchanges that take place between muscle and the blood by which it is traversed. The last method is specially applicable to warm-blooded animals ; in the first two methods the excised muscles of the frog have been exclusively employed.

To examine the gaseous exchanges between muscle and air the posterior extremities of a frog are left suspended in a closed vessel for several hours, at the end of which the air is analysed. It is found to have lost oxygen and gained carbon dioxide. We are not, however, justified in at once attributing the change to a respiratory activity of living muscle, because a control experiment made under similar conditions with dead muscle gives a similar result, the effect in this case being due to putrefaction, which, like respiration, is attended by a consumption of oxygen and a production of carbon dioxide. As regards the consumption of oxygen, it has not been found possible to distinguish a respiratory from a putrefactive effect; moreover, the consumption of oxygen bears no constant relation to the production of carbon dioxide. The oxygen item is therefore discarded from further study by this method. But as regards the production of carbon dioxide, it is possible to distinguish a true respiratory from the subsequent putrefactive discharge, and to estimate differences of the respiratory discharge coinciding with differences of muscular activity. Attention is therefore concentrated upon the carbon dioxide item.

Evidence of the above statements is supplied by the comparison of the gaseous exchanges of muscle at rest with those of tetanised muscle, as well as by the gas discharge of muscle in nitrogen or *in vacuo*. There is no demonstrable difference in the amounts of oxygen absorbed by resting and by tetanised muscle, whereas there is a marked difference in the amounts of carbon dioxide exhaled in the two cases. Fresh minced muscle subjected for many hours at a temperature of 20° to 30° to the vacuum of a mercurial pump yields a continuous discharge of gas; during a first period the discharge is considerable, it then slackens, and again becomes considerable. The first discharge is regarded as respiratory, it consists almost entirely of carbon dioxide; the second discharge is the effect of putrefaction, it consists of carbon dioxide and of nitrogen, and it continues for an indefinite period. Muscle suspended *in vacuo* as above, or in an indifferent atmosphere of nitrogen, may thus discharge carbon dioxide with no less activity than when it is in an atmosphere containing oxygen. An entire frog washed free of blood by the injection of salt solution, and enclosed in nitrogen or *in vacuo*, continues equally to discharge carbon dioxide. These facts show conclusively that carbon dioxide is not due to the immediate action of oxygen, but that it is formed in the dissociation of



some storage compound. The previous action of oxygen has no doubt contributed to the formation of any such compound; or—as Pflüger expressed it—absorbed oxygen has helped to wind up the clock, discharged carbon dioxide is the sign of its running down. To this storage substance, which, however, is not a real and identified compound, but only a hypothetical idea, Hermann has given the name ‘Inogen’ (from *ís*, *ívós*, force, or muscle), on account of the supposition that it is the force-producing material in muscular contraction.

The extraction of muscle gas, which is exclusively carbon dioxide, under various conditions of experiment, furnishes data upon which conclusions are based. By raising the temperature to 40° or 50°, muscle is made rigid or ‘rigored’; it becomes acid, and the primary discharge of gas is rapidly completed; the addition of phosphoric acid now liberates a further small quantity of carbon dioxide. The carbon dioxide removable by vacuum is termed ‘free’; that removable by acid is termed ‘fixed’: the total volume of carbon dioxide (free and fixed) thus obtainable from muscle is 1 to 15 vols. per 100. If muscle is suddenly heated to 70°, it is fixed without passing through the stage of true rigor; it is said not to acidify, and no carbon dioxide is discharged; what gas happens to be in the muscle at the time is obtainable from it by vacuum and by acid, but the hypothetical storage compound ‘inogen’ is ‘fixed’ and rendered incapable of further change. This process, technically known as ‘scalding,’ renders possible certain comparisons. Thus, comparing the amounts of gas obtainable from scalded fresh muscle and from scalded tetanised muscle, much more carbon dioxide is obtainable from the latter than from the former. Comparing scalded fresh muscle with rigored muscle, more carbon dioxide is obtainable from the latter than from the former. Rigored yields more carbon dioxide than tetanised muscle. Finally, by comparing the amounts obtainable (*a*) from fresh rigored muscle, (*b*) from muscle which was first tetanised, then rigored, Hermann found that the rigor gas was less in (*b*) than in (*a*), and that the sum of rigor and of tetanus gases in (*b*) was about equal to the rigor gas obtained from (*a*). The conclusion drawn from these observations is that one and the same substance, ‘inogen,’ suffers dissociation and yields carbon dioxide in heat-rigor and in muscular contraction.

Fresh muscle minced and boiled for two or three hours yields as much as 100 vols. carbon dioxide per 100 vols. muscle. Muscle which has been tetanised or heat-rigored and allowed to

exhale the  $\text{CO}_2$  thus produced, subsequently yields about 30 vols. per 100 on boiling. The substance from which carbon dioxide is thus produced is presumably Hermann's inogen, and we are led to suppose that, besides its physiological dissociation in muscular contraction, it dissociates slowly *post mortem* (death-rigor), rapidly at  $40^\circ$  (heat-rigor), is fixed at  $70^\circ$  (scalding), again dissociates at  $100^\circ$  (boiling). In contraction, after death, and at  $40^\circ$ , muscle yields  $\text{CO}_2$  and becomes acid; at  $70^\circ$ , although it becomes rigid, it does not yield  $\text{CO}_2$  nor become acid; at  $100^\circ$  it yields  $\text{CO}_2$  and becomes acid.

The gaseous exchanges taking place between muscle and blood have been more particularly studied in France by Bernard and by Chauveau, in Germany by Ludwig and by Pflüger. Bernard observed the fundamental fact that venous blood coming from muscles is darker during contraction than during quiescence. The difference is obviously due to the greater deoxygenating effect of active than of resting muscle, and if we take into account that the amount of blood that passes through the muscles is much greater during contraction than during rest (according to Chauveau four or five times as great), we must recognise that the absolute amount of carbon dioxide produced by muscle is much greater during contraction than during repose. Bernard's estimates do not, however, supply any exact numerical expression for this difference. Ludwig and his pupils, Schmidt and Sczelkow, attempted to fill this gap by the gas-analysis of blood made to circulate artificially through excised muscles, as well as of blood coming from living muscle *in situ*. They confirmed the fundamental fact that active muscle consumes more oxygen and produces more carbon dioxide than resting muscle, but the conditions of experiment were too far removed from the normal for it to be allowable to admit as normal the numerical data obtained. Later experiments in the same laboratory by v. Frey have shown that the consumption of  $\text{O}_2$  under such conditions is not more than  $\frac{1}{10}$ th to  $\frac{1}{5}$ th the normal, whereas the production of  $\text{CO}_2$  and of lactic acid was comparatively large. Pflüger, in conjunction with his pupils, Stintzing and Finkler, criticised and rejected Ludwig's conclusions, in particular the statement that the amount of oxygen in the blood and the amount of the latter passing through the muscle determine the energy of  $\text{CO}_2$  production. They upheld by very convincing arguments the precisely opposite doctrine, to wit, that the activity of muscle determines the oxygen consump-



tion, not that the oxygen-supply determines muscular activity. The possible connection between lactic acid and  $\text{CO}_2$  in rigor and in contraction will be considered under 'Muscle' (p. 321).

To sum up :—*The activity of muscle-respiration is determined by the activity of muscle, and not vice versa.*

*The consumption of oxygen is determined by the oxygen requirements of tissue, not by the amount of oxygen available.*

*The exhalation of carbon dioxide is the most reliable indicator of muscle-respiration.*

*By integration of oxygen a force-yielding storage substance is formed.*

*By disintegration of this substance carbon dioxide is liberated, in company with heat, work, acid, and an alteration of electrical potential.*

'Pressure' or 'tension' of gas. 'Partial pressure.'—The pressure of the atmosphere is equivalent to that of a column of mercury about 75 centimeters high. Atmospheric air is a mechanical mixture of oxygen and nitrogen—in round numbers, 1 part oxygen to 4 parts nitrogen. The pressures of oxygen and of nitrogen in the mixture are proportionate with their respective volumes, *i.e.*  $\frac{1}{5}$  the total pressure ( $=15$  cm. Hg) is the partial pressure of oxygen,  $\frac{4}{5}$  the total pressure ( $=60$  cm. Hg) is the partial pressure of nitrogen. More generally expressed, the *total* pressure or tension of a mixture of gases is equal to the sum of the *partial* pressures or tensions of the component gases. In the case of air the total pressure (75 cm. Hg) is equal to the partial pressure of oxygen (15 cm. Hg) *plus* the partial pressure of nitrogen (60 cm. Hg). Thus, if the percentage of gas in a mixture is known, the total pressure being also known, the partial pressure of the gas is obtained by a simple calculation. The percentage of oxygen in air being 20, the pressure of air being 75 cm. Hg, the partial pressure of oxygen is  $\frac{20 \times 75}{100}$  cm. Hg  $=15$  cm. Hg.

The same laws apply to simple solutions of gases in liquids when chemical forces do not intervene; but in the case of blood, in which the gases are held in loose combination, their partial pressures cannot be calculated from the percentage, and must be ascertained by direct observation. For example, in blood containing 12 vols.  $\text{O}_2$  per 100, the oxygen, in the absence of chemical force, would have a tension or partial pressure  $=\frac{12 \times 75}{100}$  cm. Hg  $=9$  cm. Hg; whereas by direct experiment its

tension is found to be much lower, *i.e.* about 2 cm. Hg. Similarly in blood containing 40 vols. CO<sub>2</sub> per 100, the CO<sub>2</sub> if simply dissolved would have a tension =  $\frac{40 \times 75}{100}$  cm. Hg = 30 cm. Hg; whereas by direct experiment its tension is found to be much lower, *i.e.* about 4 cm. Hg. The explanation in both cases is that the gases are held in loose chemical combination. Thus the blood-gases do not conform to the Henry-Dalton law of pressure, according to which the volume of gas dissolved in a fluid varies directly as the pressure; if pressure is gradually lowered in the blood-pump, the gases are not evolved *pari passu*, their dissociation from the blood at temperatures between 37° and 17° C. does not take place in abundance until the pressure is reduced to between  $\frac{1}{3}$  and  $\frac{1}{5}$  of its normal value.

The tension of CO<sub>2</sub> in the blood is ascertained by the *aëro-tonometer*. This in principle is an apparatus in which blood is brought into close relation with two gaseous mixtures, in one of which the CO<sub>2</sub> tension is above, while in the other it is below, the anticipated CO<sub>2</sub> tension of the blood. When blood is allowed to dribble through the apparatus it takes CO<sub>2</sub> from the first mixture, and yields CO<sub>2</sub> to the second, and from these data a fairly accurate estimate of CO<sub>2</sub> tension in the blood is derived. For example, if in the two parts of the tonometer the original CO<sub>2</sub> tension were equivalent to 3 and to 6 cm. Hg respectively, and if by the action of blood on the two mixtures these values were increased to 4 and diminished to 5 cm. respectively, the tension of CO<sub>2</sub> in the blood would be estimated at 4.5 cm. Hg.






The CO<sub>2</sub> tension of alveolar air is obtained by the *pulmonary catheter*; this is essentially a double tube, the outer part of which can be dilated so as to block a bronchus, while the inner channel remains in communication with the air beyond the blocked point. The bronchus being blocked, a portion of the lung is cut off from ventilation, and the tension of the contained gas soon equalises the maximum tension of gas in the pulmonary capillaries. The CO<sub>2</sub> tension in the alveolar air Wolffberg thus found to be equal to the CO<sub>2</sub> tension in venous blood, and not much greater than that of expired air.

An idea of the relative magnitudes of the O<sub>2</sub> and CO<sub>2</sub> tensions in various situations may be gathered from the subjoined table, principally composed of data supplied by the observations of Wolffberg and of Strassburg on the dog. With reference to these numbers, it is to be remarked that they denote magnitudes which are presumably lower than normally exist in the human



subject, seeing that the  $\text{CO}_2$  of expired air averages 3 per cent. ( $=2\cdot25$  cm. tension) on the dog, and 4 per cent. ( $=3$  cm. tension) on man; for blood we have taken in round numbers the tension of 3 cm. Hg as that of  $\text{CO}_2$  and of  $\text{O}_2$ , but, as might be expected, arterial and venous bloods differ in this respect—representative values are 3 cm.  $\text{O}_2$  tension, 2 cm.  $\text{CO}_2$  tension in arterial blood, 2 cm.  $\text{O}_2$  tension and 4 cm.  $\text{CO}_2$  tension in venous blood; as regards alveolar air, the value of the  $\text{CO}_2$  tension given is that of blocked alveoli; the value of  $\text{O}_2$  tension is an approximate value of what, judging from the partial pressure of  $\text{O}_2$  in the pleural cavity ( $5\cdot7$ ), presumably obtains in the unblocked alveoli; in blocked alveoli the  $\text{O}_2$  tension falls to that of the pulmonary blood, viz. 3 cm.; the difference in the  $\text{CO}_2$  tension of alveolar and expired air is not so great as might have been expected—in other words, the diffusion of  $\text{CO}_2$  is extremely rapid. As regards lymph, it is to be observed that the  $\text{CO}_2$  tension actually observed is very variable, sometimes exceeding, sometimes falling short of the  $\text{CO}_2$  tension in venous blood, and that the  $\text{O}_2$  tension is not always at zero; this is attributable to the fact that under the conditions of experiment more or less gas diffuses between lymph and the arterial blood of neighbouring vessels.

*Partial Pressure of Oxygen and of Carbon Dioxide.*

	Oxygen		Carbon dioxide
In atmospheric air . . . .	15 cm. Hg		$\cdot03$ cm. Hg
In expired air . . . . .	12 „		2·25 „
In alveolar air . . . . .	6 „		2·5 „
In blood . . . . .	3 „		3 „
In lymph . . . . .	0 „		4 „
In tissue . . . . .	0 „		6 „

The passage of oxygen from the atmosphere to the blood and to the tissues, as well as the passage of carbon dioxide from the tissues to the blood and to the atmosphere, can thus be accounted for on purely physical grounds, the incoming current of  $\text{O}_2$  and the outgoing current of  $\text{CO}_2$  being regarded as ordinary diffusion-currents. The diffusion of oxygen is directed from atmospheric to alveolar air, from alveolar air to blood, from blood to tissue. The diffusion of carbon dioxide is directed from tissue to blood, from blood to alveolar air, from alveolar to atmospheric air.

The concurrence of forces other than diffusion has been invoked to account for the discharge of  $\text{CO}_2$ . It has been shown

that the absorption of  $O_2$  is an adjuvant to the expulsion of  $CO_2$ , and also that the presence of the N in atmospheric air favours the escape of  $CO_2$ ; v. Fleischl, moreover, attributes a considerable part to the dissociating effect of the heart's contraction, which mechanically assists in the 'liberation' of  $CO_2$ ; Bohr invokes a secretory action of the pulmonary epithelium. If a comparison be made between equal volumes of blood and of serum as regards the amount and tension of  $CO_2$  in each case, it will be found that the amount is greater in serum, but that the tension is greater in blood; all the  $CO_2$  of blood is removable by the gas-pump, but in serum a certain proportion is 'fixed,' and removable only after the addition of acid. A pure solution of hæmoglobin will absorb a greater volume of  $CO_2$  than an equal amount of water, and the absorption does not conform to the Henry-Dalton law. From this it has been inferred by Bohr that  $CO_2$  forms a compound with Hb, but no such compound has yet been identified.

As regards the absorption of oxygen, there can be no doubt that the tissues themselves attract oxygen by virtue of their powerful reducing action. The pressure of oxygen in tissue is maintained at zero, the 'appetite' of tissue for oxygen is so great that not only will it admit oxygen at low pressure, but it will attract oxygen which is held back in chemical combination; thus, living tissue is capable of deoxygenating blood down to the last trace of oxygen; it can even take oxygen from compounds far more stable than oxyhæmoglobin. Of this powerful reducing action we have a striking instance in the effect of living tissue upon alizarin blue. This substance is white in a deoxidised state, blue in the oxidised state. Powerful chemical means are required to reduce it, yet living tissue in or out of the body is capable of so doing; alizarin mixed with minced muscle in a closed vessel will in a few hours lose its blue colour. This effect, be it observed, is not contradictory but complementary of the results of the gas-analysis of muscle, which only shows that the gasometric method is not suitable to the study of respiratory consumption of oxygen by muscle.

The state of tissue is the determining factor in a chain of events—active respiration of tissue increases the subtraction of oxygen from arterial blood and increases blood-flow through the muscle. Increased blood-flow is thus the consequence and not the cause of increased activity of tissue. And it may be that the acidification which accompanies muscular contraction, and



which is known to promote vaso-dilatation, constitutes one of the links between the cause and the effect.

The term 'pressure' is employed so frequently and in so many senses in connection with the mechanism of respiration, that it will be advisable to enumerate and distinguish its various applications:—1. It is used in connection with the pulmonary circulation (blood-pressure). 2. It is used in connection with the pressure of a gas mixed with other gases (partial pressure). 3. It is used in connection with the air-passages (intrapulmonary pressure). 4. It is used in connection with the pleural cavity (pleural or intrathoracic pressure). Pulmonary blood-pressure needs no special explanation; the partial pressure of gases has just been explained. Intrapulmonary and pleural pressures are frequently confused, but should be carefully distinguished from each other; the first is the pressure within the air-passages; it is measured by placing a manometer in connection with one nostril or with a T-tube fixed in the trachea, and it is positive or negative according as the chest is in expiration or in inspiration: the second is the pressure in the pleural cavity; it is measured on the living animal by placing a manometer in connection with a cannula inserted through the thorax and parietal layer of the pleura, and on the dead animal by placing a manometer in connection with a cannula tied in the trachea, and then laying the chest open. The pleural pressure is below that of the atmosphere, *i.e.* negative, in ordinary expiration, as well as in inspiration; the only case in which it becomes positive is when an expiratory effort is obstructed; it is the consequence of the elasticity of the lungs, which tend to shrink with more or less force according as the chest is more or less distended; this negative or subatmospheric pleural pressure is least in the expiratory position, greatest in the inspiratory position of the thorax; in ordinary expiration it is  $-5$  mm. Hg; in ordinary inspiration  $-10$  mm.; in maximum inspiration  $-30$  mm. The intrapulmonary pressure varies very little with normal respiration; in ordinary free expiration it is  $+2$  mm.; in ordinary free inspiration it is  $-1$  mm.; in partially obstructed expiration, as in speaking and singing, it is  $+10$  to  $+30$  mm. With completely closed air-passages, an expiratory effort may reach the pressure of  $+90$  mm., and an inspiratory effort that of  $-60$  mm. Hg.

Violent expiratory efforts, especially if made against obstruction and frequently repeated, are apt to damage the air-vesicles;

the pressure within them may be made so great that they become permanently over-distended and may even be ruptured, giving rise to a condition which is termed 'emphysema.' The lungs of persons who blow wind instruments, and of patients subject to frequent and prolonged fits of coughing, are particularly liable to become thus altered.

**Influence of respiration upon circulation.**—Respiration affects circulation in three distinct ways; the movements of respiration have a direct mechanical effect upon blood-pressure, they also influence the heart and the blood-vessels through nervous channels, and, thirdly, the more or less oxygenated state of the blood,

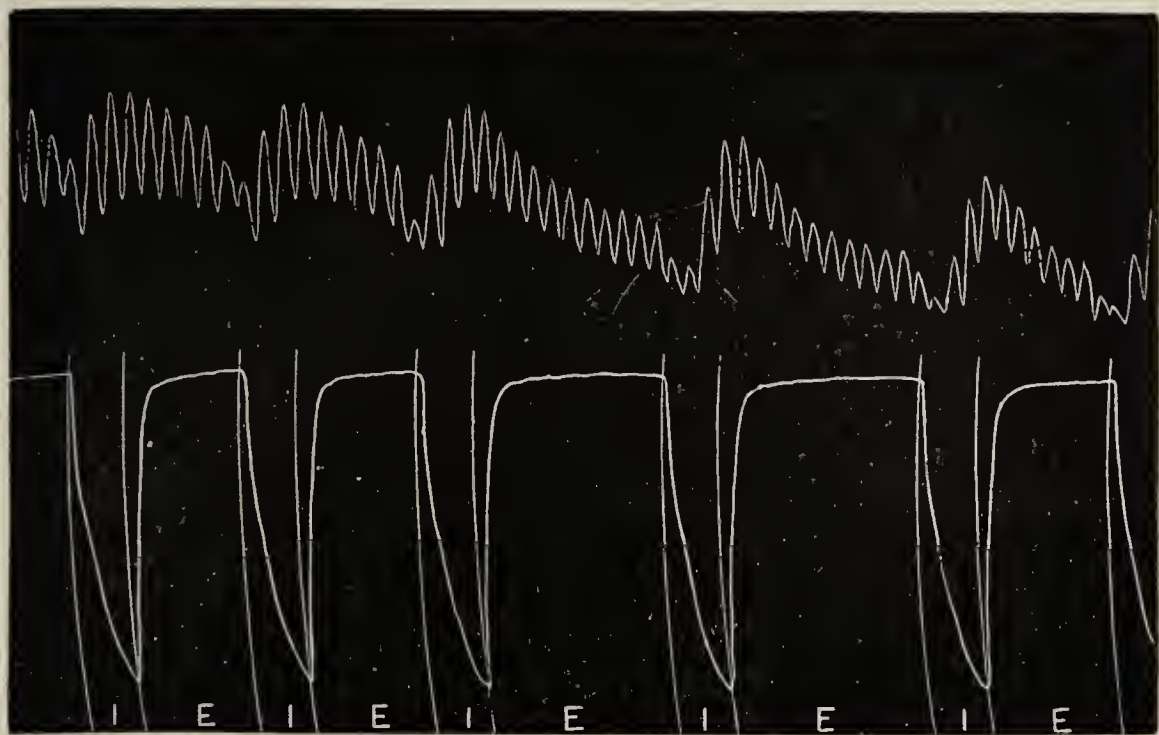


FIG. 63.—RABBIT. INFLUENCE OF RESPIRATORY MOVEMENTS UPON ARTERIAL BLOOD-PRESSURE.

brought about by respiration, takes effect upon the vasomotor centre.

Every blood-pressure tracing shows the effect of the respiratory movements; if these be simultaneously recorded with the blood-pressure, it will be found that the arterial pressure arises with inspiration and falls with expiration—the inspiratory rise beginning a little later than the beginning of inspiration, and the expiratory fall beginning a little later than the beginning of expiration. These are mechanical effects of the varying intra-thoracic pressure, as will presently be explained.

The clearest evidence that respiration influences the circulation through nervous channels, modifying the beat of the heart, is furnished by the dog; on this animal the pulse-frequency is considerably greater during inspiration than during expiration, a difference which is attributable to the vagi nerves, since it is abolished by their section.



The influence of respiration on the arterial tone is likewise transmitted through efferent nervous channels, and is attributable to the quality and quantity of the blood-supply to the spinal bulb. When the blood is deficient in oxygen, the bulb is stimulated, the vasomotor centre and nerves are excited, and the blood-pressure rises: when the blood-supply of the bulb is obstructed, as by compression of the carotids, the same effects ensue; they are equally well produced on an animal with both vagi divided and kept alive by artificial respiration. The excitation of the bulbar vasomotor centre by a deficiency of oxygenated blood cannot endure for any long period; the excitability of its grey matter is soon exhausted if oxygenated blood is permanently withheld: when this occurs the arterioles begin to relax and the blood-pressure falls to and below the normal. This fall, which may last one or two minutes, is not uniform and continuous, but characterised by undulations of pressure, due to rhythmic vaso-constrictor efforts, the last gasps of the dying vasomotor centre. That these undulations (generally known as Traube-Hering curves) are certainly due to vasomotor action, and are not mechanical effects of respiratory efforts, is proved by the fact that they occur in the absence of respiratory movements; they are, moreover, of slower rhythm, not more than two or three per minute, and thereby distinguishable from the mechanical effects of respiration, even while these are taking place.

As regards an explanation of the mechanical influence of respiratory movements upon blood-pressure, it is necessary to refer to the mechanical conditions to which the heart and vessels are subject in the thoracic cavity. If the thorax were a freely open cavity, its expansion and contraction could not exercise any pressure on the heart and large blood-vessels; air would simply pass in and out of the cavity. If the thorax were a rigidly closed cavity, *i.e.* if the trachea were closed, each expansion would diminish the intracardiac pressure, *i.e.* exercise a suction action upon the heart and large vessels; each contraction would increase the intracardiac pressure, *i.e.* exercise a squeezing action upon the heart and large vessels. The thorax may for the moment be regarded as a cavity with a limited outlet, so that its contraction and expansion exercise a slight and temporary plus and minus pressure in the air-passages during expiration and inspiration.

Moreover—and this is the most important factor of the

normal respiratory undulations of blood-pressure—the lungs are elastic, tending to shrink away from the thoracic walls, so that everything in the thorax outside the lungs—viz. pleural cavity, heart and intrathoracic vessels—is at a more or less subatmospheric pressure, according as the lung exercises more or less elasticity. Or, otherwise regarded, the heart and intrathoracic vessels are not under the full atmospheric pressure of 760 mm. Hg; the air-column does not bear directly upon them, but only indirectly through the distended lung, which by virtue of its elasticity tends to shrink, thereby supporting a certain proportion of atmospheric pressure; the actual air-pressure upon the heart and intrathoracic vessels is therefore subatmospheric, viz. 760 mm. Hg, *minus* the elastic force exercised by the distended lung. The value of this force is about 5 mm. Hg in the position of ordinary expiration, and about 10 mm. in the position of ordinary inspiration. So that the heart and intrathoracic vessels exercise a suction of 5 to 10 mm. upon the contents of extrathoracic vessels in the two phases of normal respiration. Taking into account the intrapulmonary pressure of +2 during expiration, and -1 during inspiration, the total suction exercised will be -3 in expiration and -11 in inspiration.

The thick-walled ventricles and the aorta, in which the blood-pressure is high, will be least influenced by this extravascular variation of pressure, which does not amount to 10 mm. of mercury. The right auricle and the venæ cavæ, which are thin-walled and at almost zero pressure, will on the other hand be sensibly affected by such a variation of pressure, which will practically determine the amount of blood-flow to the right side of the heart. Increased or diminished blood-flow to the right side of the heart rapidly entails increased or diminished blood-flow to the left side, which therefore drives forward a larger quantity of blood soon after inspiration has begun, a smaller quantity of blood soon after expiration has begun. Arterial pressure goes up in the first case, goes down in the second.

The pump-action of the expanded lung as just described is no doubt the most effective agent in furthering the flow of blood from the systemic veins to the right auricle, and consequently furnishing the blood which is driven on by the left ventricle; certain special cases have yet to be considered.

*In laboured or obstructed respiration* the chief varying factor



is the exaggerated intrapulmonary pressure, which of course entails a corresponding alteration of the pleural pressure. A forcible inspiration entails a greater negative pressure in the air-passages and in the pleural cavity, a forcible expiration entails a greater positive pressure in the air-passages, and the pleural pressure is now positive instead of negative. These effects are well marked when the respiration is partially obstructed or exaggerated—the great veins at the root of the neck shrink and swell visibly with inspiration and with expiration; the effects are at a maximum when respiratory efforts are made during closure of the mouth and nose—under these circumstances the total inspiratory suction of  $-11$  mm. may be increased to  $-70$ , and the expiratory suction action of  $-3$  mm. may be replaced by a repulsive action amounting to  $+90$  mm. Hg. A violent and prolonged inspiratory effort with closed mouth and nose can even cause a temporary arrest of the circulation, the intrathoracic vessels being distended with blood and the auricles unable to contract; this experiment, which is not without discomfort or even danger, is known as Müller's experiment. Conversely, a violent and prolonged expiratory effort with closed air-passages is capable of arresting the circulation; the intrathoracic vessels are in this case emptied of blood, and the venous blood from the system is prevented from entering the compressed right auricle; this is known as Valsalva's experiment.

Breathing *compressed or rarefied air* influences the blood-pressure, as might be anticipated from the above considerations. Breathing in compressed air lowers the blood-pressure, breathing in rarefied air raises the blood-pressure. If arrangements be made by means of valves to inspire from rarefied and to expire into compressed air, the normal inspiratory increase and expiratory decrease of blood-pressure will be exaggerated; if, on the contrary, inspiration be taken from compressed air, and expiration be made into rarefied air, the normal variations may be diminished, abolished, or even reversed. A man on the top of a mountain breathes rarefied air, in a diving-bell he breathes compressed air, the endurable range of pressure being from a minimum of  $\frac{1}{2}$  an atmosphere to a maximum of 5 atmospheres, which are the pressures obtained at an altitude of 5,500 meters above sea-level and at a depth of 40 meters under water. Death occurred in a balloon ascent reaching 8,600 meters—an altitude at which the pressure is only

$\frac{1}{3}$  atmosphere.<sup>1</sup> At a pressure of 15 atmospheres animals die in convulsions, at 20 atmospheres germination and putrefaction are arrested, *i.e.* no living cell can breathe. Sudden variations of pressure are far more dangerous than gradual alterations; the pressure to which a diver is submitted within his diving-dress must be very gradually lowered when he is raised to the surface; there is at this time danger of accidents (*e.g.* paraplegia) due to embolism by a disengagement of gases within the blood.

The *vera causa* of 'mountain sickness,' of which the most prominent symptom is dyspnœa, is a diminished atmospheric pressure; persons who undertake the exertion of a climb without preliminary training are particularly liable to an attack of this nature; seasoned mountaineers even are apt to become affected at an altitude of between 16,000 and 17,000 feet.

In *artificial respiration*, as ordinarily performed by bellows, the intrapulmonary pressure is increased with inflation, diminished with collapse; the pressure-conditions are thus the reverse of those obtaining in normal respiration, and the blood-pressure undulations (provided a moderately slow respiratory rhythm be chosen) are reversed. Each inflation (=inspiration) causes an immediate short rise, followed by a more prolonged fall of blood-pressure; each extraction (=expiration) causes an immediate fall, followed by a more prolonged rise. These effects are attributable to alterations of air-pressure in the pulmonary alveoli; inflation first squeezes the blood on towards the heart and then retards further blood-flow by compressing the pulmonary capillaries; extraction relaxes the intra-alveolar pressure, it thus momentarily limits the blood-flow and subsequently permits it in greater volume.

Alterations of this kind, which are undoubtedly effective in artificial respiration as above described, are probably of some account even in normal respiration; inspiration relaxing the bed, and causing a slight preliminary retardation of blood-flow through the lung, followed by acceleration; expiration compressing the bed, and causing a preliminary acceleration, followed by retardation. But this is at most a very accessory factor in the production of the normal undulations, which have been fully accounted for as the consequence of the pumping action of the lungs. Another accessory factor is perhaps of some moment, *viz.* the

<sup>1</sup> According to presumably trustworthy reports, the greatest height attained in balloon ascents was 8,838 meters, and the greatest depth by pearl-divers 54 meters below water.



compression of the abdominal viscera, more especially of the liver, by the inspiratory descent of the diaphragm; kneading or 'massage' of the abdomen is a ready and effectual means of promoting venous flow, thus raising arterial pressure, and it is possible that ordinary respiration has some action of this kind.

The influence of the respiratory movements upon venous pressure has been incidentally alluded to in the foregoing description, and is easily realised; the familiar fact that the veins shrink with inspiration and swell with expiration is of itself sufficient to remind us of the general rule that variations of venous pressure are in a contrary sense to variations of arterial pressure. This rule holds good throughout all the cases above considered; we may therefore briefly summarise the main effects observed in the venous and arterial pressure—*e.g.* of a carotid artery and of a jugular vein—in the following form:—

			Car. Press.	Jug. Press.
Normal respiration	{	Inspiration . .	+	—
		Expiration . .	—	+
Artificial respiration	{	Inflation . .	—	+
		Extraction . .	+	—

**Control of respiration by the nervous system.**—The movements of normal respiration are automatic and involuntary, but subject to occasional modification by the stimulation of afferent nerves and by voluntary interference. The nerve machinery consists of (1) a chief respiratory centre in the spinal bulb, (2) afferent nerves, (3) efferent nerves to the muscles of respiration. In its normal automatic mechanism the respiratory centre acts in obedience (1) to the chemical state of the blood as regards the amount of gases, and (2) to the mechanical state of the lung as regards distension. In occasional modifications to which the mechanism is subject the centre acts in response (1) to exaggerations of its habitual stimuli, (2) to voluntary mandates, and (3) to centripetal stimuli along almost every afferent nerve.

*The respiratory centre.*—Destruction of the spinal bulb at once arrests the movements of respiration; and provided the bulb be left intact, destruction of the brain does not abolish these movements. This, of itself, is enough to show that the spinal bulb includes the chief respiratory centre, even though it is not possible to define the centre anatomically as this or that nucleus of grey matter. The most we are able to say with

any approach to certainty is that this centre is above the vaso-motor centre, and that each lateral half of the bulb is principally concerned in the control of respiratory movements upon the same side of the body.

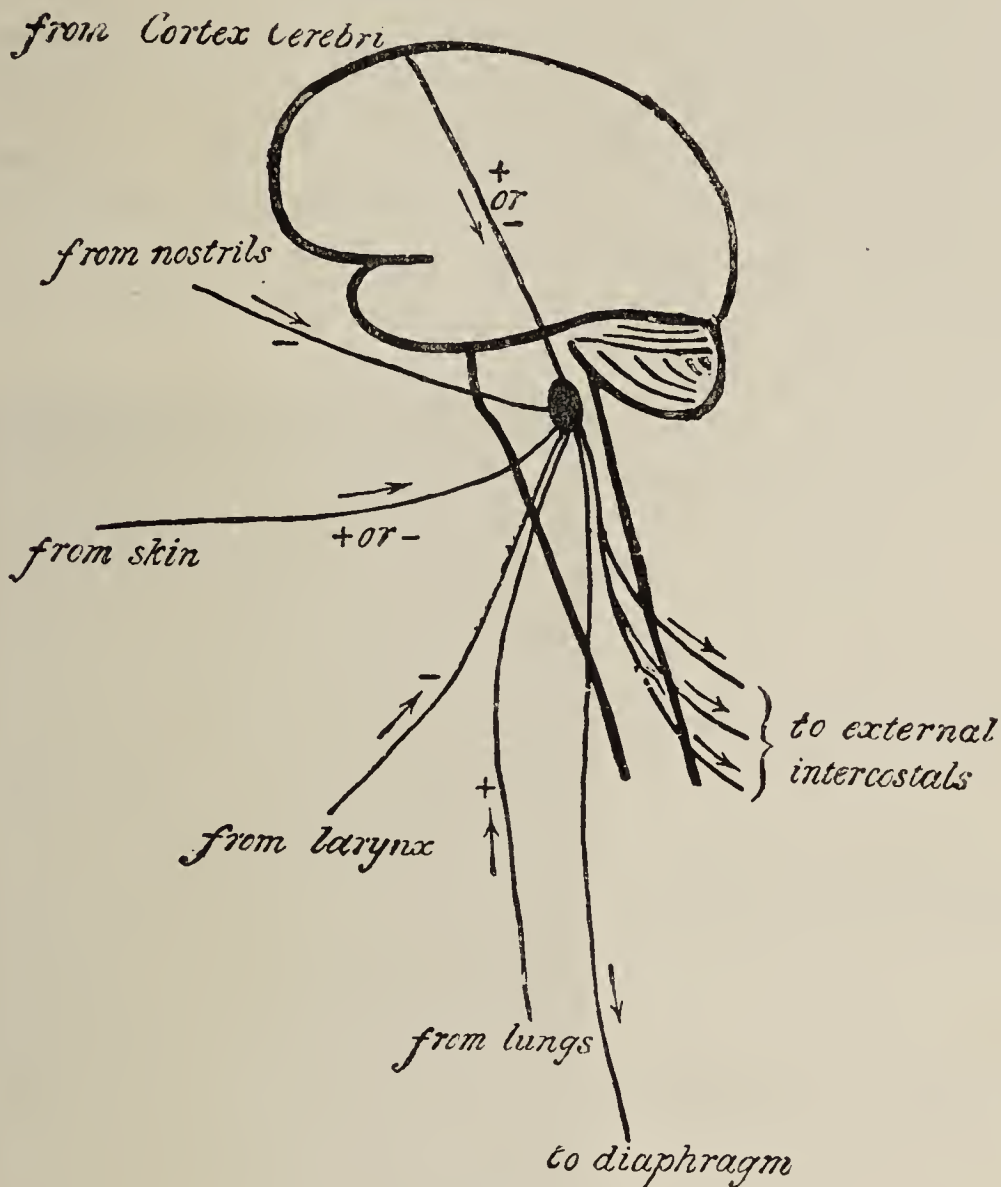


FIG. 64.

Diagram to illustrate the chief nervous connections of the respiratory centre. Impulses may reach it—

- (1) From the cortex of the brain (*voluntary control, emotional modifications*).
- (2) From the surface of the body (*increase or diminution by cutaneous stimuli*).
- (3) From the lung by way of the vagus (*increase*).
- (4) From the larynx by way of the superior laryngeal (*diminution*).
- (5) From the nostrils by way of the fifth nerve (*diminution*).

Impulses pass from the centre to the diaphragm along the phrenic nerve, to the intercostal muscles along intercostal nerves, and in laboured respiration along many other nerves to many other muscles.

The expression 'chief respiratory centre' implies a certain reservation. Observations on animals poisoned with strychnine, more especially on young animals, have been recorded which prove that imperfect respiratory movements persist after destruction of the bulb; these movements are accepted as evidence that the upper part of the spinal cord, in a minor and subordinate



degree, takes part in the control of respiration. But this share is less considerable than in the case of vasomotor action, which, as we have seen, while chiefly controlled from the bulb, is also to some extent controlled from the spinal cord.

The respiratory centre is particularly sensitive to the quality of the blood by which it is traversed ; its excitability is instantly raised by blood which is deficient in oxygen ; it is, on the contrary, lowered by blood which is fully saturated with oxygen, and in consequence of these alterations of excitability the centre increases or diminishes the energy of respiratory movement. Thus the respiratory supply is a self-adjusting process ; ill-oxygenated

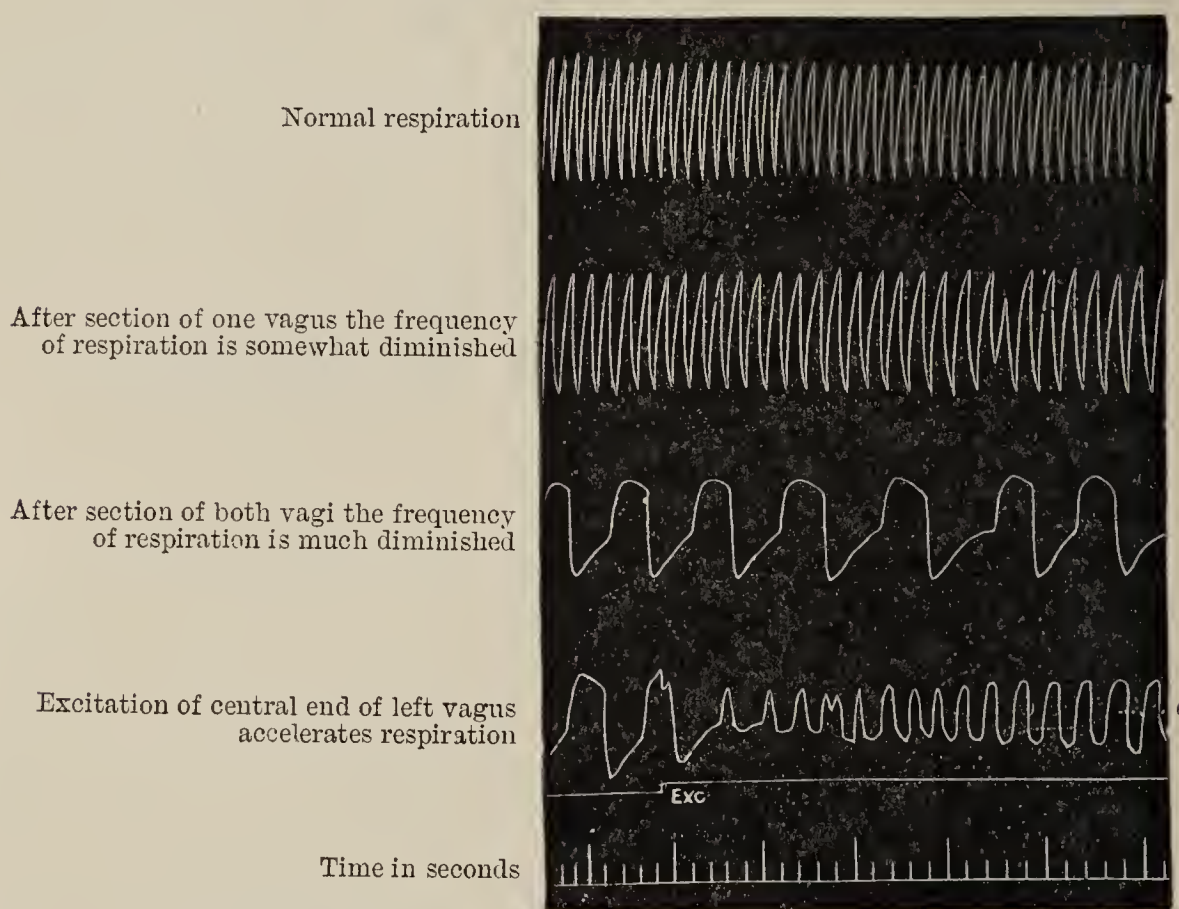


FIG. 65.—INFLUENCE OF THE VAGUS UPON RESPIRATORY MOVEMENTS.

blood excites respiration and thereby makes up arrears, well-oxygenated blood calms respiration and thereby cuts down excess, the respiratory centre in this relation playing the part of 'blood-taster' to the whole body.

The afferent nerves in most intimate relation with respiratory acts are, in order of importance, the vagus, the superior laryngeal, and the fifth nerve. The vagus appears to be in constant action ; the effect of section of both vagi is to render the respiration very slow ; excitation of the *central* end of either vagus, if moderate, quickens the rhythm up to and beyond the normal frequency ; if excessive, it quickens the respiration to such an extent as to arrest respiration in inspiratory tetanus ; at the same time the

glottis is found to open wide. These facts show that the vagi are afferent channels of a *plus* influence from the lung that augments respiration. This is the rule, but it is liable to exceptions; one vagus, more often the left than the right, may be the more effective; excitation of the central end of a vagus may sometimes diminish the frequency of respiration instead of increasing it, and lead to arrest in expiration instead of in inspiration; moreover the nature of the excitation is not a matter of indifference; according to Langendorf an unfailing demonstration of inspiratory augmentation may be made on a chloralised rabbit by exciting the central end of the vagus with a *descending* current of one or two cells interrupted ten times per second; while an uninterrupted *ascending* current of the same strength gives expiratory arrest. It has also been pointed out by Knoll that the mere fact of raising the nerve from its bed and letting it down again may be sufficient to excite an expiratory pause; the excitation in this case is probably a sudden variation of 'nerve current,' and the observation shows what a delicate matter this vagus effect must be, and what extreme care is required before positive conclusions may be drawn. The effects may also differ according to the part of the nerve to which excitation is applied; this last circumstance is, however, intelligible when the effects of stimulating its superior laryngeal branch have been learned. Section of one or of both these branches has no effect, but excitation of the central end of either, if moderate, slows the rhythm; if excessive, arrests the chest in respiration. We learn from this that the superior laryngeal may upon occasion transmit a *minus* influence from the larynx which diminishes respiration, and we may, indeed, recognise it to be natural that irritation of the larynx by a foreign body should suspend the inspiratory, and, it may be, promote the expiratory, act. Very similar effects follow excitation of the nostrils, from which the nasal branch of the fifth is the sensory nerve; respiration is diminished, and we may recognise this also as an appropriate effect. These are the three nerves excitation of which gives the most marked results, but the excitation of any afferent nerve may alter respiration, only we cannot always predict whether it will increase or whether it will diminish the act. A dash of cold water to the surface of the body may stop respiration or may start it, so may painful stimulation of the central end of a sensory nerve; but the result may not safely be predicted; the same nerve can 'blow hot and cold'; at most we may expect that weak stimuli will more probably



accelerate respiration, strong stimuli more probably arrest respiration.

The more or less distended state of the lung is an important factor in the alterations of respiratory movements, by giving rise to stimuli which reach the bulb *viâ* the vagus nerve; distension is unfavourable to inspiration and favourable to expiration, collapse is favourable to inspiration and unfavourable to expiration; thus each phase of respiration tends to limit itself and to promote its opposite by the stretching and shrinking of the pulmonary alveoli, or, otherwise expressed, inspiration inhibits inspiration and excites expiration, expiration inhibits expiration and excites inspiration (Hering).

Up to this point we have considered the part played by the vagus as an afferent nerve; to conclude our account of the relation of the vagus to the lung, it should be mentioned that by virtue of efferent fibres this nerve exercises motor action upon the bronchial muscle, *i.e.* is a broncho-constrictor nerve. This action has long been assumed to exist, more especially in the theory of spasmodic asthma, but until quite recently its physiological

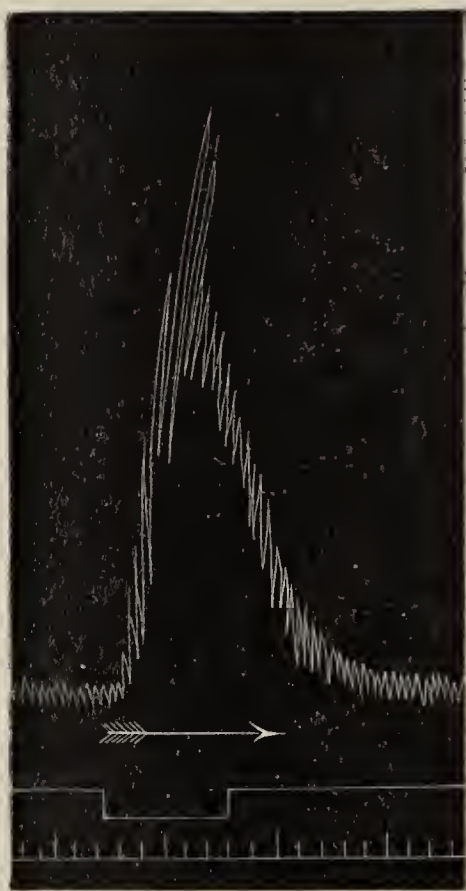


FIG. 66.

Effect of vagus excitation on intrapulmonary pressure as shown by the rheotome method. (Einthoven.)

demonstration has been very incomplete; it has, indeed, been shown by means of a manometer tied into the trachea that the intrapulmonary pressure is slightly raised by excitation of the vagus, but the broncho-constrictor action thus indicated cannot be brought into full evidence. A complete demonstration of broncho-constriction has been given as follows: a curarised dog is submitted to artificial respiration, the inflations being made at regular intervals and with uniform quantities of air; a side-tube from the tracheal tube leads to a manometer, by way of a special tap, which is automatically opened by means of the respiration apparatus for a definite short period of each inflation, thus letting a short

puff of pressure into the manometer. If the resistance to inflation is constant, the succession of puffs produces a level

series of manometer effects ; but if the bronchial muscle be made to contract by excitation of the peripheral end of the vagus, the puffs encounter greater resistance, and the volume of inflation being unaltered, they must produce a greater effect upon the manometer. A constriction of minute bronchioles, which might cause little or no effect upon a *steady* intrapulmonary volume or pressure of air, will produce a very evident effect when the rising and falling phases are, so to speak, 'tapped' on this 'window plan,' which is similar in principle to the method of leading off an electrical pressure by the rheotome (p. 312).

Einthoven has thus found: (1) that the excitation of the peripheral end of either vagus causes strong broncho-constriction, shown by increase of pressure in and distension of the lungs; (2) that this action is abolished by nicotine or by atropine; (3) that broncho-constriction is caused by inhalation of  $\text{CO}_2$ . The broncho-constrictor effect of vagus excitation begins in about 1.2 to 1.3 seconds, and wears off very rapidly during excitation; the effect on the manometer may amount to 175 mm.  $\text{H}_2\text{O}$  or more.

### THE LARYNX

The lungs form part of the organ of voice, constituting as they do a bellows supplying the blast of air which, in consequence of obstructions and resonations in the larynx and in the mouth, furnishes the vibrations of musical tones and of articulate sounds. The voice is produced by the vibrations of the vocal cords influenced by the expiratory blast from the lungs; the articulate sounds of speech are produced by the interruptions and resonations of voice in the laryngeal, pharyngeal, and buccal cavities caused by the tongue and lips. In man and in many animals — dog, sheep, horse, &c.—the voice is usually an expiratory act; other animals, such as the cat, pig, and ox, vocalise with inspiration.

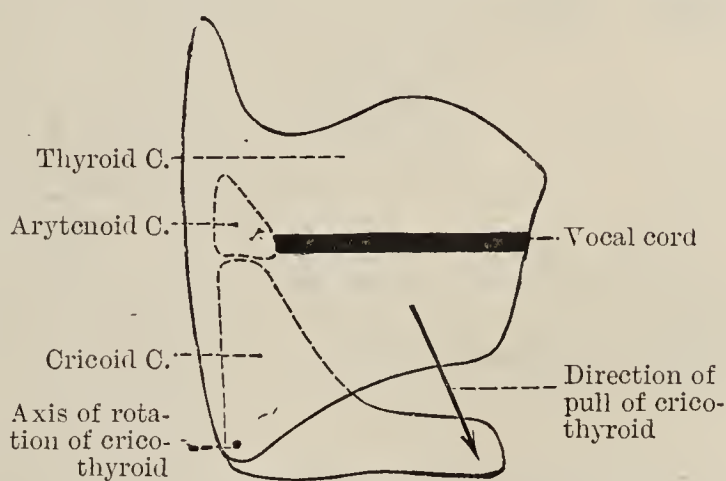


FIG. 67.—DIAGRAMMATIC VERTICAL SECTION OF LARYNX.

The *pitch* of sounds and tones yielded by the larynx depends upon the tension of the vocal cords; with a high-pitched tone



the vocal cords are extended, with a low-pitched tone they are shorter and comparatively relaxed. The quality or timbre of the voice is chiefly influenced by resonations in the mouth, nose, and accessory air-cavities.

The larynx consists of a framework of *cartilages* united by *ligaments* and moved by six pairs of *muscles*. The cartilages are the thyroid, cricoid, and two arytenoids; the most important ligaments are the thyro-arytenoid ligaments or true vocal cords; the six pairs of muscles are the crico-thyroid, the lateral crico-arytenoid, the posterior crico-arytenoid, the thyro-arytenoid, the inter-arytenoid, and the aryteno-epiglottidean. The *vocal cords* stretch across the top of the larynx from the vocal process of the

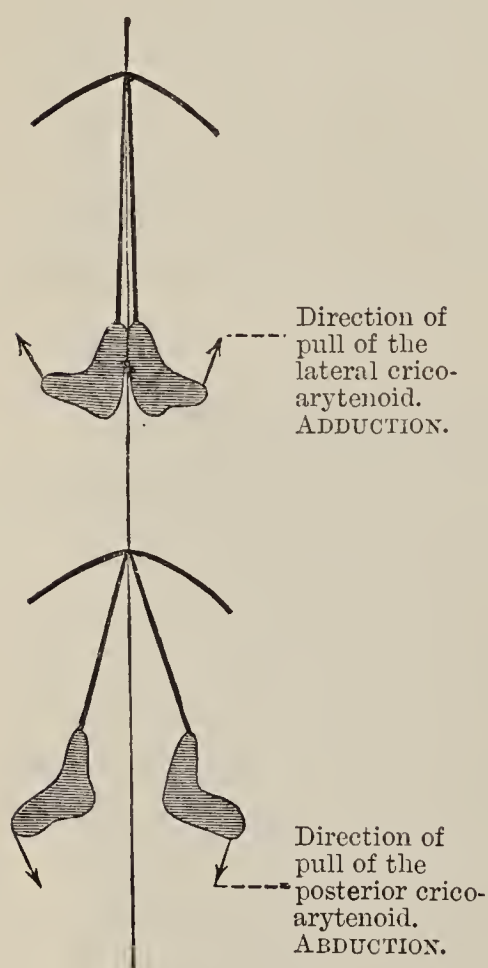


FIG. 68.

Diagrammatic horizontal section through the arytenoid cartilages and vocal cords to illustrate the action of the crico-arytenoid muscles.

arytenoid cartilage to the angle of the thyroid rather below the salient point that is seen externally as the *pomum Adami*. The length of the relaxed vocal cords averages 18 mm. in the male and 12 mm. in the female, and the space or chink between their free edges is the glottis or rima glottidis. The articulation between the thyroid and cricoid cartilages is such that contraction of the crico-thyroid muscle will tilt the posterior part of the cricoid backwards, or the anterior part of the thyroid forwards, thus stretching and elongating the vocal cords. The articulation between the cricoid and arytenoid is such that the principal movements of an arytenoid cartilage are movements of rotation round a vertical axis, carrying the vocal process and vocal cord towards or from the middle line: these movements of adduction and abduction are effected by the muscles at-

tached to the muscular process of the arytenoid cartilage, viz. the lateral and the posterior crico-arytenoid. These two pairs of muscles are in fact the most important of the laryngeal muscles; the lateral crico-arytenoids rotate the arytenoid cartilages inwards and adduct the vocal cords; they are the muscles most concerned in vocalisation; the posterior crico-

arytenoids rotate the arytenoid cartilages outwards and abduct the vocal cords; they act in association with the muscles of inspiration, widening the aperture of the glottis during inspiratory movements. The inter-arytenoid and aryteno-epiglottidean muscles draw the arytenoid cartilages together and thus effect closure of that portion of the glottis which otherwise remains patent between the two cartilages. The thyro-arytenoid muscles (divisible into an external and an internal portion) have the same general direction as the vocal cords, and in part lie between and in actual attachment to their upper and lower folds; they draw the arytenoid cartilages towards the thyroid, thus relaxing the vocal cords; by their internal portions they are capable of tightening an anterior while relaxing a posterior part of the

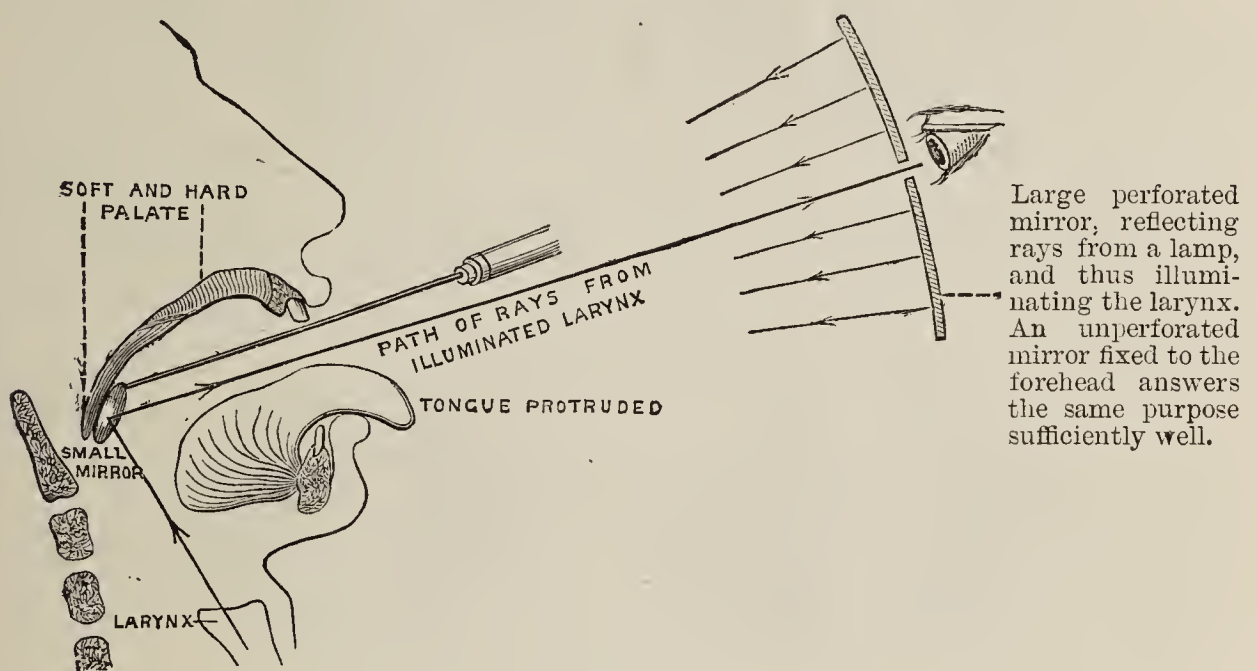


FIG. 69.

To illustrate the manner in which a view of the interior of the larynx is obtained.

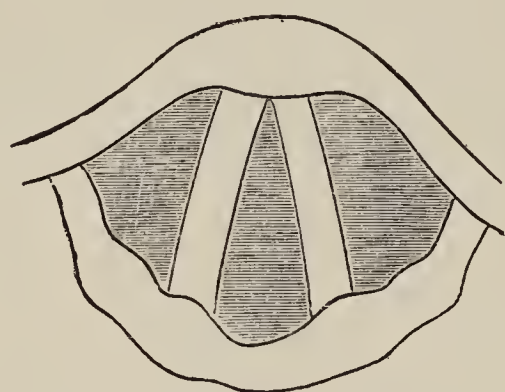
cords, and it is possible that they may exercise a 'stop' action upon the cords, causing them to vibrate in partial lengths instead of in their whole length.

The tension of the vocal cords can also be modified by external muscles attached to the thyroid-cartilages, more especially the sterno-thyroid and thyro-hyoid; the former depresses the thyroid cartilage and tilts it backwards, thus relaxing the vocal cords; the latter raises the thyroid and tilts it forwards, thus stretching the vocal cords. The inferior constrictor of the pharynx may also assist in relaxing the tension of the vocal cords of a sufficiently flexible larynx by flattening the thyroid cartilage.

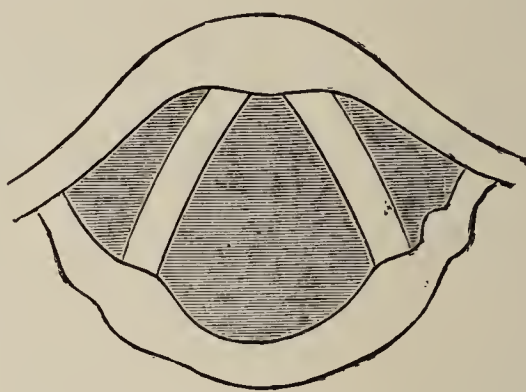
The interior of the larynx and the movements of the vocal cords are visible on man and on animals by means of the *laryngo-*



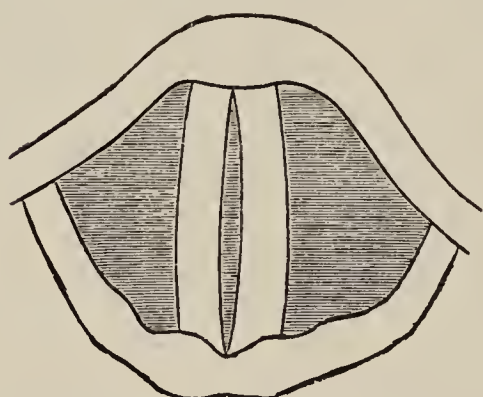
*scope*. This is a contrivance consisting essentially of two mirrors, a larger perforated mirror fixed in front of the eye of the observer, a smaller mirror at the end of a handle by which it can be held at the back of the mouth; the observer looks through the perforated mirror by means of which he reflects the light of a lamp in the same direction, and inspects a reflection of the interior of the larynx from the smaller mirror held at a suitable angle. The image seen is reversed, the apparent left cord is in reality the right cord, the arytenoid ends of the cords appear



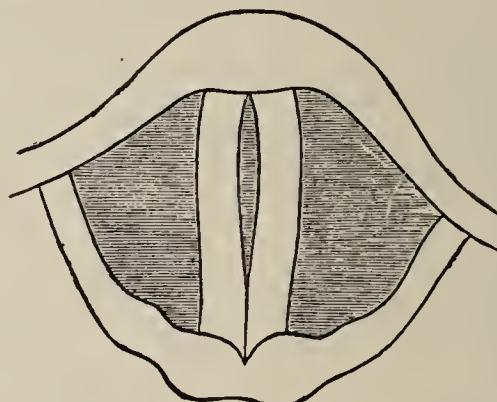
Position of rest. (The vocal cords are midway between abduction and adduction.)



Position during forced inspiration. (The vocal cords are in extreme abduction.)



Position during vocalisation. ('Chest voice.') The vocal cords are adducted and vibrating in their entire length.



Position during vocalisation. (Falsetto voice.) The vocal cords are adducted and vibrating in their anterior portions only.

FIG. 70.

Diagrams to illustrate the position of the vocal cords under various circumstances.

anterior, &c. Thus viewed, the vocal cords of a normal larynx are seen to move away from each other during inspiration, and to return towards the middle line during expiration; during the utterance of a sound the cords are seen to come quite close together, the glottis being reduced to a mere chink.

With regard to the position of the cords and consequent shape of the glottis during the singing of notes in different registers, considerable differences of opinion have prevailed, more especially as regards falsetto or head notes. The accompanying figures illustrate the more generally credited view, that the cords

vibrate in their whole length in the natural emission of tones, but that they are 'stopped' (by the action of the internal part of the thyro-arytenoid) and only vibrate by their anterior portions during the emission of a falsetto tone. According to other observers a more or less deep margin of the edges is set into vibration in the different registers, the vibrating margin being narrowest in falsetto.

At moments of extreme muscular exertion, when the chest is filled, and fixed with an expiratory effort, the larynx is tightly closed; in this closure the false as well as the true vocal cords take part. In a cough as excited by contact of a foreign body with the mucous membrane of the larynx, the glottis is closed during the first part of a sudden expiratory effort, and suddenly gives way to the expiratory rush of air.

The *nervous supply* of the larynx is derived from the superior and from the inferior laryngeal branches of the vagus. The inferior laryngeal is the chief motor nerve of the laryngeal muscles, supplying the lateral and posterior crico-arytenoids, the thyro- and inter-arytenoids, and in part the crico-thyroid. The superior laryngeal is the sensory nerve of the laryngeal mucous membrane; it also takes part in the motor supply of the crico-thyroid.

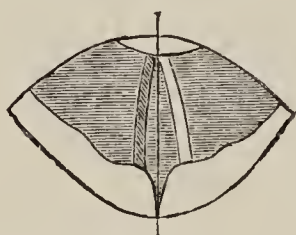
The consequences of paralysis or of excitation of the laryngeal nerves are in some respects simple, in others peculiar. Clinically as well as experimentally, the commonest form of laryngeal paralysis is caused by section or by compression of the inferior laryngeal nerve; if the nerve of one side be cut or paralysed—say on the right side—the voice is lost (aphonia) because the right vocal cord cannot be adducted, breathing is impeded (inspiratory dyspnoea) because it cannot be abducted; on laryngoscopic examination the right vocal cord neither moves from the middle line with inspiration, nor towards the middle line with vocalisation. It is interesting to note that in a gradually produced paralysis of the inferior laryngeal it is—contrary to expectation—the abductor or less voluntary muscle, the posterior crico-arytenoid, which fails first and degenerates most, the adductor or more voluntary muscle, the lateral crico-arytenoid, which fails at a later stage; but this order of failure is reversed when function is abolished by an excessive administration of ether.

Electrical stimulation of the peripheral end of the inferior laryngeal nerve, calling into play all the laryngeal muscles,

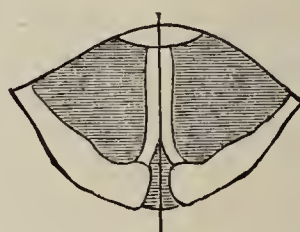


abductors, as well as adductors, produces effects that vary with the conditions of experiment. Ordinarily, *i.e.* upon an adult dog, and with moderately strong stimulation of a fresh nerve the adductors predominate, producing closure of the glottis; but a reverse effect, *viz.* abduction, is not uncommon, *e.g.* upon puppies, or with weak stimulation, or during deep etherisation.

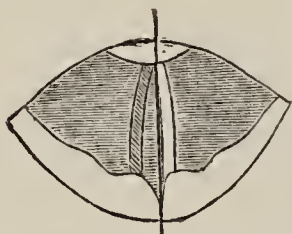
Paralysis or section of the superior laryngeal nerve abolishes the sensibility of the larynx, and it consequently fails to resent the intrusion of foreign bodies, which under normal conditions are expelled by coughing. The voice is rendered hoarse in consequence of deficient action of the crico-thyroid and deficient



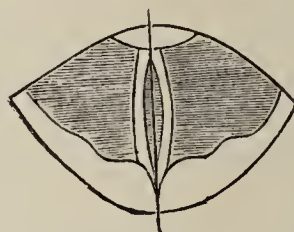
Paralysis of right abductor or of right inferior laryngeal nerve. Position during inspiration. The right vocal cord fails to be abducted, and remains nearer the middle line than the normal left cord.



Paralysis of inter-arytenoid. Position during vocalisation. The arytenoid cartilages fail to come together, the inter-cartilaginous portion of the glottis remains unclosed.



Paralysis of right adductor, or of right inferior laryngeal nerve. Position during vocalisation. The right vocal cord fails to be adducted, and remains further from the middle line than the normal left cord.



Paralysis of thyro-arytenoid. Position during vocalisation. The vocal cords are not properly stretched, and their edges remain unduly curved.

FIG. 71.

Diagrams to illustrate the chief paralytic defects in the movements of the vocal cords (mirror views).

tension of the vocal cords. Excitation of the central end of the divided nerve produces reflex movements, of deglutition, reflex arrest of respiratory movements, and closure of the glottis.

Paralysis of individual muscles produces effects that can easily be deduced from their known physiological action. Thus paralysis of adductors interferes with the expiratory adduction necessary to vocalisation, and causes loss of voice; paralysis of abductors interferes with the inspiratory abduction necessary to free breathing, and causes inspiratory dyspnoea: paralysis of the tensors (crico-thyroid, internal portion of thyro-arytenoid) and of the inter-arytenoids causes hoarseness, in the first case

because the vocal cords cannot be properly stretched and approximated, in the second case because the arytenoid cartilages cannot be brought together (see fig. 71).

Paralysis of one side of the brain (hemiplegia) does not affect the laryngeal muscles on either side, the movements of the vocal cords are unimpaired; on the other hand, excitation of one side of the brain sets in action the laryngeal muscles of both sides, the vocal cords are adducted (usually) or abducted (sometimes).

The laryngeal and other nerves concerned in the movements of speech have their spinal centres of origin in the bulb; bulbar paralysis causes, among other consequences, impairment of vocal pronunciation.



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**The general drift of digestion.**—‘*Food*’ includes the following classes of proximate principles—*proteids*, *carbohydrates*, *fats*, *salts*, and *water*. Of these, the last two—viz. salts and water—can pass unaltered into the animal body by simple *diffusion* through any part of the mucous membrane; they do not require to be digested. Food belonging to either of the first three classes cannot pass unchanged; it requires to be altered before it can pass into the body. This necessary change is, properly speaking, *digestion*, and it is carried on at the surface of the digestive tract, more particularly in the stomach and in the small intestine. We recognise, then, that digestion is a preparatory process by which food in the digestive tract, though not yet in the body, is altered and rendered capable of being absorbed, and we commonly include in the process this, its last step, viz. *absorption*, or the actual passage of the digested food through the epithelium of the mucous membrane into the underlying chyle and blood-vessels. Before absorption, food is outside the body, though in the intestinal tube undergoing digestion: after absorption, it is in the body forming part of the nutritive fluids, blood and lymph. We may, indeed, by analogy with the process of respiration, speak of digestion as an external process—a transaction between the contents of the intestine on one side and the blood and lymph on the other. Food in the intestine is acted upon by the series of secretions poured out by the glands in the walls of the digestive tract, and by glands (the pancreas and the liver), the ducts of which open into that tract. These secretions, which are derived from the blood, are:—saliva, gastric juice, bile, pancreatic juice, and intestinal juice, and their respective shares in the entire process are as follows:—Proteids are digested by the agency of gastric and of pancreatic juice; fats are digested by the agency of bile and of



pancreatic juice ; starch, by the agency of saliva and of pancreatic juice ; cane-sugar by the agency of intestinal juice. In all these cases (except in that of cane-sugar) the effect of digestion is to transform indiffusible into diffusible bodies, and thus facilitate their absorption. But that absorption is not physically identical with diffusion is illustrated by the case of cane-sugar, which, although highly diffusible, does not pass unchanged, and by that of fat, which in part at least can pass, although not strictly speaking diffusible. We recognise, then, that the absorption of digested matter depends upon two conditions : firstly, a physical condition—the diffusibility of the digested material ; secondly, a physiological condition—the selective activity of the epithelium through which absorption is effected. The essential constituent of each digestive fluid is a *ferment*—*ptyalin* in saliva, *pepsin* in gastric juice, *trypsin* in pancreatic juice, *invertin* in intestinal juice. These and other ferments are the specific agents by which the digestive transformations of food are effected. They are the occult agents of modern physiology, inasmuch as they have never been isolated as definite bodies, and are recognised to be present only by the effects they produce. They are not confined to animals, but are also found in plants.<sup>1</sup>

The chemical action of the digestive fluids is promoted by mechanical assistance ; each mouthful of food is subdivided by *mastication* and propelled into the stomach by *deglutition* ; the swallowed mass is churned and well mixed with gastric juice by the muscular action of the stomach-wall ; reduced to pulp (‘chyme’) it is squeezed through the pylorus and propelled onwards along the intestinal tube by *peristaltic contraction* ; finally, what is left of it—the refuse which has not been absorbed—together with a small proportion of actual excretion from the intestinal tube, accumulates in the rectum, and is discharged by the movement of defæcation.

**Physiological anatomy of the digestive tube.**—The digestive tube is lined internally by mucous membrane, externally by muscle, and receives the ducts of several tributary glands (salivary, hepatic and pancreatic). The mucous membrane is the channel of absorption and of secretion, the muscular coat supplies motor power

<sup>1</sup> Insectivorous plants (*Drosera* ; *Dionæa*) contain peptic ferment. Tryptic ferment has been identified in the fruit and leaves of *Carica Papaya*, rennet ferment in the common bedstraw (*Galium verum*) and in many other plants, lipolytic ferment in the castor-oil plant (*Ricinus communis*) ; diastatic ferment—converting starch into sugar—is widely distributed, and is most readily obtained from germinating barley (malt).

by which solid food is broken up and propelled along the canal. In the greater part of its length, and excepting more particularly at its two extremities (mouth and anus), the muscular coat of the tube is composed of involuntary smooth fibre disposed in two layers, internal circular and external longitudinal, between which there is a close plexus of non-medullated nerve-fibres and ganglion-cells, the myenteric or Auerbach's plexus. The mucous membrane differs in different segments; from œsophagus to rectum it has a layer of smooth muscle (*muscularis mucosæ*) as one of its constituents, and it is separated from the muscular coat proper by a submucous connective-tissue coat, in which are contained the ramifications of blood-vessels, lymphatics, and nerves, the latter forming a loose plexus of non-medullated fibres and ganglion-cells (the plexus of Meissner). In the mouth and in the œsophagus the epithelium of the mucosa is stratified, in the stomach it is beset with branched tubular glands, in the small intestine with simple short tubular glands (Lieberkühn's) and with projections (villi), in the large intestine with long tubular glands; at the anus stratified epithelium again appears. The mucous and submucous coats contain, moreover, a large quantity of lymphoid tissue, especially at the tonsils, in the lower half of the small intestine (Peyer's patches), and in the large intestine. The greater part of the abdominal portion of the tube is covered externally by a serous coat formed by the peritoneum. The terms 'inner' and 'outer' are used with reference to the tube itself; the tube is, however, an involution of the external surface, so that physiologically, with reference to the body-mass, the epithelium of the mucous membrane is the true external surface.

The mucous membrane of the tongue is beset with variously shaped projections or papillæ, anatomically distinguished as filiform, fungiform, and circumvallate. The latter, confined to the posterior part of the tongue, are each surrounded by a circular trench, on the sides of which the so-called '*taste-buds*' are found. These are small ovoid bodies composed of long cells, to which nerve-fibres have been traced; they are supposed to be gustatory in function, although the fact has not actually been proved. The substance of the tongue is striated muscle, the fibres of which are of small diameter and branched. The muscles of the lips, cheeks, palate and pharynx are of the ordinary skeletal character.

The teeth are composed of dentine, surrounding a cavity which contains the 'pulp'; the exposed portion or crown of the



tooth is covered by enamel, the embedded part or root by cement ; vessels and nerves and lymphatics ramify in the pulp, the mass of which consists of lymphoid tissue.

### THE DIGESTIVE MOVEMENTS

**Mastication.**—Each mouthful of solid food requires to be reduced to a more or less minutely subdivided state by chewing. This state is of great importance to subsequent digestion by facilitating the free access and admixture of the digestive juices with the food ; considerable lumps of food may indeed be disposed of by persons having a ‘good digestion,’ but in weaker, and especially in old persons, a hurriedly swallowed meal is apt to be followed by indigestion, and bad teeth are a fruitful source of dyspepsia, *i.e.* difficult digestion, simply owing to the fact that imperfectly chewed food is digested with difficulty.

Each person possesses during life two complete sets of natural teeth—a first or temporary set, the milk-teeth, appearing during the first two or three years of life, and being displaced by a second or permanent set, which begin to make their appearance about the sixth year. The milk-teeth are twenty in number, and comprise eight incisors or cutting teeth, four canines or tearing teeth, eight molars or grinding teeth. The permanent teeth are thirty-two in number, and comprise eight incisors, four canines, eight pre-molars or bicuspid, and twelve molars. A comparison of the skull and teeth of man with those of a purely carnivorous animal, such as the tiger, and with those of a purely graminivorous animal such as the ox, leads to the conclusion that a mixed diet comprising flesh and vegetables is natural to man. A comparison of their intestines leads to the same conclusion ; the intestine of a flesh-feeder is short, that of a vegetable-feeder is long, that of a man is of moderate length.

**Deglutition.**—A mouthful of food, having been masticated and mixed with saliva, is collected by the muscles of the lips, tongue, and cheeks into a rounded mass or ‘bolus,’ which glides over the base of the tongue into the pharynx, where it is grasped and passed onwards by the action of the constrictor muscles to the œsophagus, along which it is forced into the stomach by a succession of muscular contractions. The act of deglutition is thus divisible into three stages—1, *buccal* ; 2, *pharyngeal* ; 3, *œsophageal*. Of these, the first, or buccal stage is entirely voluntary ; so long as the bolus has not passed through the isthmus of the fauces it

may be rejected if desired ; once past the isthmus it cannot be turned back.

The second or pharyngeal stage is an involuntary and highly complicated reflex act, composed of several distinct movements, which concur to close the superior orifices of the pharynx, and prevent the food or liquid from returning to the mouth or penetrating to the nose or larynx. This result is effected by (1) contraction of the pharynx, the muscles of which grasp the bolus ; (2) contraction of the isthmus of the fauces, shutting off the buccal from the pharyngeal cavity ; (3) elevation of the larynx and occlusion of the glottis ; (4) elevation of the velum palati and occlusion of the posterior nares, shutting off the nasal from the pharyngeal cavity. The shape and position of the epiglottis are such as to suggest that it might act as a lid covering the superior orifice of the larynx during the second stage of deglutition ; but it may be lost without loss of protection, and it is chiefly because the larynx is raised and the base of the tongue thrust backwards that food and liquids do not normally find access to the wind-pipe. The state of the orifices of the Eustachian tubes during the second stage of deglutition has been debated ; it is probable that these orifices are closed during inaction of the pharynx, and opened during the second stage of deglutition ; hence the usual practice of inflating the tympanum by forcing air through the nostrils while the patient is in the act of swallowing.

The third or œsophageal stage completes the act of deglutition. The contraction, which is an involuntary one, is of an apparently 'peristaltic' character ; that is to say, it is a slowly progressing wave of constriction. It is remarkable, however, that the wave is not arrested by section or ligature of the œsophagus, whereas it is arrested by section of the œsophageal nerves (Mosso). These facts show that the apparently peristaltic wave is the expression of a sequence of action occurring in the nervous centre with which the œsophageal nerves are in functional connection, and not a true local wave of contraction along a continuous sheet of muscle. Stimulation by the passage of swallowed food or by saliva is required to produce the contraction, which is therefore reflex in character. When experimentally excited the contraction can only travel from above downwards, and not in the reverse direction ; but the phenomena of rumination and vomiting are significant of the possibility of a reversed order of progression when contraction occurs under peculiar physiological conditions.



*Time relations.*—The time relations of a complete act of deglutition are on man as follows. Taking the contraction of the mylo-hyoid muscle as the point of departure,

the contraction of the pharynx commences				0·3 sec.	later
that of the upper part of the œsophagus				1·2	„
„	middle	„	„	3·0	„
„	lower	„	„	6·0	„

and (corresponding with the fact that on man the upper part of the œsophageal muscle is striped, the lower part smooth, and the middle part mixed) the duration of the contraction is in the first part 2 to 2·5 seconds, in the second part 6 to 7 seconds, in the third part 9 to 10 seconds ; thus the contraction outlasts the passage of food, which takes about 6 seconds. In the dog's œsophagus, which is of striped muscle throughout, the contraction has in all parts a duration of between 1 and 2 seconds.

*Sounds caused by deglutition.*—On auscultation of the œsophagus of man during the deglutition of fluid, two distinct sounds are audible—the first practically simultaneous with the commencement of the act, the second about 6 seconds later. The first (which is not always to be heard) is attributable to a squirt of liquid down the œsophagus ; the second (which can nearly always be heard) is probably due to the peristaltic contraction of the œsophagus squeezing the fluid through the cardiac orifice of the stomach. Kronecker and Meltzer have shown that if a series of voluntary acts of deglutition are made at intervals of a second, it is not until the last of these that a ‘peristaltic’ contraction of the food-tube is produced. Each act of deglutition interferes with the natural consequence of the preceding act which should have led to a complete contraction, and it is probable that the glosso-pharyngeal nerve is the afferent channel of this peculiar form of arrest. In the passive state the cardiac end of the œsophagus is semi-contracted ; it relaxes during the last part of deglutition, and contracts more firmly after the conclusion of the act.

*Nervous mechanism.*—Deglutition, after completion of the buccal stage, is a well-characterised and definite reflex act. It is not possible to swallow without something to swallow ; and it is the contact of food or of liquid with the anterior surface of the soft palate which supplies the necessary stimulus, the afferent channel being the fifth nerve ; it may also be accidentally as well as experimentally provoked by excitation of the mucous membrane of the larynx, or of the superior laryngeal branch of the

vagus. A third channel through which the reflex act of deglutition is influenced, but in an opposite sense, is the glosso-pharyngeal nerve; an act of deglutition experimentally excited by stimulation of the central end of the superior laryngeal, or by irritation of the soft palate, fails to take place during excitation of the central end of the glosso-pharyngeal, thus showing that the last-named nerve exercises a centripetal inhibitory effect on the stimulus. The centre from which the movements of deglutition are governed is situated in the upper part of the spinal bulb; destruction of the brain above, or abolition of its function, does not interfere with deglutition; on the other hand, difficulty of deglutition is one of the symptoms of disease of the medulla oblongata ('bulbar paralysis'). The efferent channels are the motor nerves to the various muscles concerned—viz. the hypoglossal (to the tongue), the motor portion of the fifth (to the mylo-hyoid), the glosso-pharyngeal, vagus and accessory (to the muscles of the pharynx, œsophagus, and palate). As has been already mentioned, the orderly contraction of the muscular tube from throat to stomach is regulated in the bulbar centre, not being interrupted by section of the œsophagus.

An experiment by Goltz is often quoted to show that the vagi nerves counteract tonic contraction of the œsophagus. The experiment is as follows:—Two frogs are hung up, one with the vagi intact, the other with both vagi cut or simply pithed. The hearts of both are removed. Salt solution is poured into their mouths. The œsophagus and stomach of the first frog are soon distended with fluid, those of the pithed frog remain obstinately contracted. In other words, a viscus from which nervous influence is removed remains in tonic contraction, while a viscus which is still under control of the nervous system yields to the pressure of fluid. In mammals, however, the vagus (by fibres derived from the spinal accessory) acts as a motor nerve upon the pharynx and œsophagus; if it is paralysed, deglutition is interfered with, and food accumulates in the œsophagus.

**Gastric and intestinal movements.**—The movements of the stomach and of the intestines are undoubtedly peristaltic in nature. Those of the stomach are of a complex churning character; those of the small intestine afford the typical example of peristaltic contraction, *i.e.* a prolonged contraction slowly advancing along the muscular tube, in the form of an annular constriction travelling along the contractile substance as a wave



which is directly transmitted from part to part, independently of excitation through nerves. The peristaltic wave of the small intestine is thus fundamentally different from the so-called peristaltic wave in the œsophagus ; the former requires only muscular continuity, and will persist after severance from the central nervous system ; the latter, as stated above, persists after section, but fails after its nerves have been severed ; moreover, the œsophageal contraction caused by local stimulation can only travel downwards, while the intestinal wave can travel in either direction, although normally it always travels downwards.

*Influence of nerves.*—The activity of intestinal movements, although not directly dependent upon, may be influenced by, nerves ; it is also modified by the state of the blood, by the nature of the chyme, and by drugs. The nerves that influence peri-

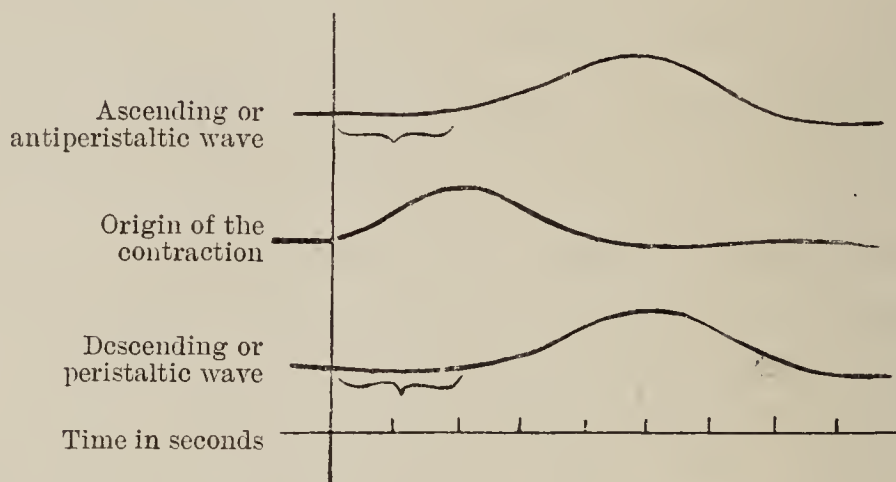


FIG. 72.

Wave of peristaltic and of antiperistaltic contraction in small intestine of dog. After Engelmann. The records were taken 5 cm. above and below the origin of the contraction ; the wave travelled this distance up or down in about 2 sec., *i.e.* at the rate of 25 mm. per second.

staltic movements are the vagus and the abdominal splanchnics. Section of any or of all these nerves causes no very obvious effect, intestinal movements continuing as before, being, if anything, more active in consequence of the vascular engorgement which is produced by section of the nerves. If, while the intestines are in active movement, the peripheral end of the splanchnic be excited the movements will be arrested. If, while the intestines are quiescent, the peripheral end of the vagus be excited, movements will be provoked ; but it must be added that the same effect is not seldom produced by excitation of the splanchnic. These results, although not so clear and decisive as might be wished, are nevertheless admitted as evidence that the vagus excites, while the splanchnics inhibit, intestinal movements. It is to be observed

that in experiments of this nature the intestines must not be freely exposed to excitation by cold air, but should be protected by immersing the animal in a warm bath of normal saline.

So long as the intestine is properly supplied with oxygenated blood, it remains comparatively quiescent; as soon as the oxygen supply is interfered with, the intestines begin to move. This interruption of the oxygen-supply can be caused in different ways, by dyspnoea with resultant venosity of the blood, by sluggish or failing circulation and consequent venous engorgement of the visceral blood-vessels, by anæmia of the intestine in consequence of the vaso-constrictor action of the splanchnic nerves. Any of these causes may excite the intestines to exaggerated movements, which are thus among the signs of failing circulation—in a fainting fit, or at the approach of death.

In the normal state of health, intestinal movements are quietly effected without arousing consciousness; but if the intestinal contents be abnormal in quality or consistence, the intestine may move too weakly or too strongly and make its presence felt. This is one among other proofs that the intestinal nerves contain afferent fibres.

Sluggish intestinal movements may be stimulated by purgative drugs, excessive intestinal movements may be controlled by sedative drugs. The consideration of these matters belongs to practical medicine rather than to experimental physiology, and it will suffice to mention, in this place, drugs the action of which has been most studied by experiments involving direct observation of intestinal movements, viz. *morphia* and *bella-donna*, which exercise a direct sedative action; *muscarin*, *colocynth*, and purgative salts, which on the contrary excite intestinal movements as well as intestinal secretion.

*Defæcation* is a reflex act, which can be initiated or resisted by voluntary action. The normal excitant of the act is the presence of *fæces* in the rectum; the normal consequence of the excitant is the orderly sequence of movements termed *defæcation*. These movements comprise:

1. Contraction of the rectum and of the abdominal muscles.
2. Contraction of the levator ani and of the sphincters.

A distended rectum is an excited and contracting rectum, it causes in consciousness a desire to *defæcate*. The voluntary response to this suggestion may be 'yes' or 'no'; 'yes' taking the form of contraction of the abdominal muscles with a passive condition of the sphincters, or 'no,' contraction of the



sphincters with a passive condition of the abdominal muscles. In this latter case the desire is resisted, and the contracting rectum drives back its contents into the sigmoid flexure. In the former case, the desire is satisfied by a normal act of relief, which commences with contraction of the rectum and of the abdominal muscles, and ends by contraction of the levator and sphincter ani. The foregoing analysis applies to the normal state, in which the excitant is of moderate degree, and when it is possible to choose between the two alternatives. It may happen, however—(1) that the excitant is excessive, the desire imperative, the voluntary negative insufficient to arrest the act: such a condition is termed incontinence of fæces or diarrhoea; (2) that the excitant is deficient, the desire absent, the voluntary affirmative insufficient to commence the act: such a condition is termed ‘constipation.’

The rectum and the anal muscles are supplied by the hypogastric and erigentes nerves; in the dog these nerves take origin from the spinal cord on a level with the 5th, 6th, and 7th lumbar vertebræ, to which region the term *ano-spinal centre* has accordingly been applied. The nervous mechanism of defæcation is closely analogous with that of micturition. (See p. 240.)

*Vomiting* is a border-land phenomenon between the normal and the abnormal events occurring in the living body. In ruminating animals the return of food from the stomach to the mouth is a regular event; in infants the act of vomiting follows the most trifling causes, and is effected with the greatest ease; in adults it is an accident that may be caused by irritation of the stomach, or, indeed, by irritation of any viscus; whereas the horse and the rabbit, in consequence of the anatomical disposition of the cardiac orifice, cannot vomit. As regards its nervous mechanism, the act itself is a reflex; its afferent channels are the afferent nerves from the seat of irritation (palate, fauces, stomach, uterus, intestine, liver, kidney, brain); its efferent nerves are the motor nerves of the stomach, of the diaphragm, and of the abdominal muscles; its centre—the so-called vomiting centre—is situated in the spinal bulb. As regards the muscular mechanism of the act, it is effected by simultaneous contraction of the abdominal muscles and of the stomach; the pyloric orifice is firmly closed, the cardiac end of the œsophagus is relaxed, and the contracting stomach is at the same time compressed by the diaphragm and by the abdominal muscles; its contents are thus driven up the œsophagus and ejected through the mouth or nose.

By experiments on animals it has been shown that the contraction of the stomach alone, or of the abdominal muscles alone, is sufficient to accomplish this ejection, for it can be completed by the stomach alone after section of the phrenic and intercostal nerves, or by the abdomen alone after the stomach has been excised and replaced by a bladder. In man, as has probably been experienced by most people, vomiting is preceded by a feeling of nausea, by a copious flow of saliva, and by preliminary 'retching' efforts, which have the effect of opening out the food-tube by distending it with air.

Drugs that cause vomiting are called *emetics*; their action may be either local, by excitation of the pharynx, œsophagus, and stomach, or general, by excitation of the spinal bulb through the medium of the blood, into which an emetic has been absorbed. The popular remedy, mustard-and-water, is an instance of an emetic acting by local excitation; apomorphine, on the other hand, is an emetic by its action on the bulb after it has been absorbed or injected into the circulation.

## THE DIGESTIVE FLUIDS

### I. *Saliva*

**Its composition.**—Ordinary saliva in the mouth is a mixture composed of the secretions of the parotid, submaxillary, sublingual, and small buccal glands. The average daily secretion amounts to  $1\frac{1}{2}$  litre. Its reaction is alkaline. It has a specific gravity of about 1,005. It is composed of water, salts, mucin, serum-albumin, and serum-globulin; it contains a ferment (ptyalin), and in solution a gas (carbon dioxide). Serum-albumin and serum-globulin are principally contained in the parotid secretion, mucin in the submaxillary and sublingual secretions; ptyalin exists in the secretions of all these glands in man, but in animals the fermentative effects on starch of different kinds of saliva are very variable. Thus, the parotid secretion is very active in the case of the rabbit (contrasting with the submaxillary of the same animal, which has little or no action), quite inactive in that of the horse, and only slightly active in that of the dog.

The chief salts present in saliva are sodium chloride, calcium carbonate and phosphate (the two latter salts are apt to form concretions as 'tartar' on the teeth, and 'salivary calculi'), and frequently, but not always, sulphocyanide of potassium; it is remarkable that so poisonous a salt should be found, even though



only in minute quantity, in normal saliva; a red coloration on the addition of ferric chloride is the test of its presence. The only gas present in quantity in saliva is carbon dioxide—100 c.c. of saliva containing as much as 60 c.c. of  $\text{CO}_2$ : only minute amounts of oxygen and of nitrogen are present, not sensibly more than in ordinary water.

A drop of saliva examined under the microscope is seen to contain a few squamous epithelial cells (which are derived from the mucous membrane of the mouth), 'salivary corpuscles,' and perhaps fragments of food and bacteria. None of these things are specially characteristic of saliva; they are only accidental impurities; the 'salivary corpuscles' themselves are simply escaped leucocytes, probably derived from the lymphoid tissue of the tonsils and of other parts.

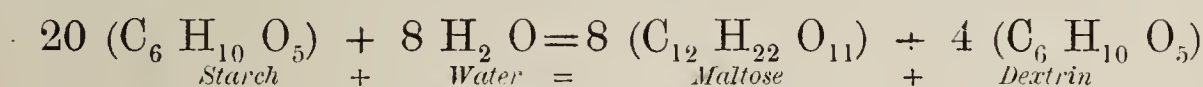
*Its uses and properties.*—The uses of saliva are mainly mechanical, and in small degree chemical. Saliva is necessary to adequate mastication and to deglutition; neither of these acts can be effectually performed with the mouth dry; the fluid is equally necessary to adequate articulation, and the sense of taste depends upon saliva to this extent, that before any solid substance can be tasted, it must be in part dissolved, saliva acting as the solvent if no fluid is taken into the mouth. The chemical action of saliva is the conversion of *starch* into *sugar* (maltose), and this conversion is brought about by the salivary ferment, *ptyalin*. That the conversion is effected by a ferment is at once proved by comparing the action of boiled with that of unboiled saliva; boiled saliva does not convert starch into sugar. The conversion by normal saliva is not immediate, nor is it complete. It is not immediate—starch acted upon by saliva passes through a series of *dextrins* before sugar is produced; among these are distinguished *erythrodextrin* and *achroodextrin*. It is not complete—the sugar produced does not reach an amount corresponding to the original amount of starch, however long the action is allowed to continue.

*Experiment.*—A mixture of soluble starch and saliva is allowed to digest at a temperature of  $35^\circ$  to  $40^\circ$ , and tested from time to time. Soluble starch gives a blue colour with iodine solution, which disappears on heating and returns on cooling. The digest of starch and saliva soon loses this reaction, and a sample of the mixture gives a red colour, which likewise disappears on heating and returns on cooling; this red colour indicates the presence of erythrodextrin. A little later the same test gives no colour

while the test for sugar does not yet give evidence of the presence of sugar; at this stage dextrin is present giving no colour with iodine, viz. achroodextrin. Later still sugar is formed in greater and greater abundance; a sample of the mixture boiled with a drop or two of copper sulphate and excess of caustic potash gives a red or yellow precipitate of copper suboxide, indicative of the presence of a reducing sugar; at this period sugar (maltose) and dextrin (achroodextrin) are present, and, however long the digestion be continued, dextrin continues to be present; the sugar produced is a reducing sugar, most nearly resembling, though probably not identical with, maltose. We have in the above experiment distinguished the following stages:—

		Tests
1. <i>Initial substance</i>	Soluble starch	Blue with iodine.
2. } <i>Intermediate sub-</i>	{ Dextrins (including erythrodextrin)	Red with iodine.
3. } <i>stances</i>		
	{ Dextrins (achroodextrin)	{ No reaction with iodine. No sugar reaction.
4. <i>Final product</i>	Dextrin and sugar	{ Yellow ppt. of $\text{Cu}_2\text{O}$ on boiling with $\text{CuSO}_4$ and $\text{KHO}$ .

The chemical equation which most nearly represents the whole process is



i.e. the process is one of ‘hydration’ or assumption of water by starch under the influence of the ferment.

The character of action of the salivary ferment is further defined by experiments showing—1, that it is destroyed by boiling; 2, that its action is delayed or suspended at a low temperature, most pronounced at about body temperature ( $37^\circ$ ); 3, that it acts best in a neutral or in a faintly alkaline medium, not at all in an acid medium, or in too strong an alkaline medium; 4, that it has almost indefinite power, if the product of its action (sugar) is not suffered to accumulate. In all these respects, with the exception of the third, the salivary ferment resembles ferments in general, which are destroyed by heat, delayed by cold, and are limited in their action only by the accumulated product of such action.

**Structure of salivary glands.**—All the salivary glands are of the compound racemose type, their branching ducts terminate—or rather commence—in groups of short tubes, the *acini*, which



are lined by cubical or spheroidal epithelium. Two chief types of gland are distinguished—(a) *serous*, of which the parotid is the representative, yielding a watery secretion with serum-albumin as its principal constituent, (b) *mucous*, of which the dog's submaxillary is the chief representative, discharging a more viscid secretion containing mucin. As usually seen in stained sections the two varieties are recognised by the greater size of the acini and of the cells by which they are lined, and by the clearness of these cells in mucous glands, as compared with the smaller acini, smaller and more granular cells obscurely separated from each other, that are characteristic of serous glands. Another noteworthy point of distinction is the presence in mucous glands of the so-called *crescents* or *demicircles*; these are groups of young cells crowded in between the fully-developed mucous cells and the basement membrane upon which they are set; they stain deeply with logwood or carmine. With regard to the termination of nerves in the gland-cells, while we cannot doubt from the results of experiments that nerves do communicate with the secreting cells of the alveoli, yet we are constrained to admit that all attempts at microscopic demonstration of the junction have been inconclusive.

Glands of the same name have by no means the same structure in all animals, nor is a salivary gland necessarily entirely 'mucous' or entirely 'serous'; it may be 'mixed,' *i.e.* serous in some parts, mucous in others; and even a single alveolus may contain individual serous and mucous cells side by side. The submaxillary of man is an instance of such a mixed gland, whereas in the dog and cat it is entirely mucous, and in the rabbit entirely serous. The parotid does not, however, seem to vary from its regular type, which is that of a serous gland in all animals.

*Changes of structure consequent upon activity.*—The microscopic appearances of the salivary glands differ greatly according as they have been at rest or in activity just before their removal. In both serous and in mucous glands alike, the acini and the cells are shrunk in the active as compared with the rested gland; the active gland has obviously expended some of its substance, as indeed is evident by comparing the entire gland of the fatigued side with the gland of the opposite side, which has been left at rest; the former is the smaller and lighter of the two.

To appreciate in detail the microscopic differences between these two contrasted states, it is necessary to bear in mind what

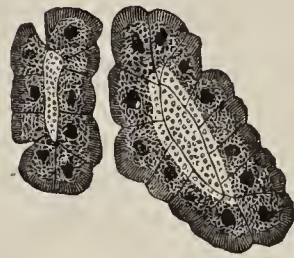
kind of preparation for microscopical examination has been adopted, and whether a given description refers (a) to alcohol-hardened gland stained by logwood or by carmine, or (b) to gland examined during life or freshly teased-out in salt solution.

Comparing carmine-alcohol preparations of a serous gland (such as the pancreas of the dog), it is observable that the cells



RESTING.

Outer stainable zone (=protoplasm).  
Inner unstainable zone (=protoplasm-product, 'zymogen').



AFTER ACTION.

Increase of outer zone.  
Decrease of inner zone.

FIG. 73.—SEROUS GLAND. DOG'S PANCREAS; ALCOHOL-CARMINE PREPARATION. (Heidenhain.)

lining the acini of the resting gland are marked by two distinct zones:—(1) an outer stained all but homogeneous zone, (2) an inner unstained and granular zone; after prolonged action the second zone is much diminished, the first zone is increased, and at the same time the whole acinus and its individual cells are considerably smaller. It is held that the outer stain-



RESTING.

Outer clear zone (protoplasm).  
Inner granular zone (zymogen).



ACTIVE.

Increase of outer zone.  
Decrease of inner zone (discharge of zymogen).

FIG. 74.—SEROUS GLAND. RABBIT'S PANCREAS OBSERVED DURING LIFE. (Kühne and S. Lea.)

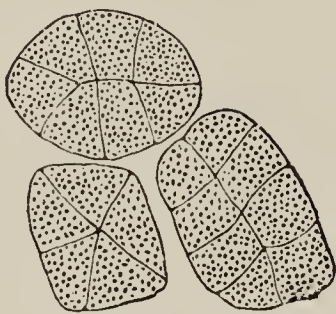
able zone contains protoplasm, while the inner non-stainable zone contains the protoplasm-product about to be secreted, and some authorities moreover hold that the visible granules of this zone are the actual 'zymogen' or precursor of ferment, which in the act of secretion becomes ferment.

The microscopical changes above described demonstrate



the discharge of material that certainly occurs in secretion, and, according to Heidenhain, they are demonstrative of a double process—(a) discharge of protoplasm-product, as shown by diminution of the granular unstainable zone, and (b) renewal of protoplasm, as shown by increase of the homogeneous stainable zone.

It has been found possible to actually observe during life changes taking place in the pancreas, that are absolutely demonstrative of the discharge of visible material from secreting cells; in the rabbit the pancreas is scattered between the layers of the mesentery, so that individual lobules may be examined by high powers of the microscope. Examined thus, a lobule is seen to be composed of a narrow outer zone which is homogeneous, and of a broader central zone which is granular; Kühne and Lea succeeded in observing, that with the process of secretion



RESTING.  
Uniform granular cells.



AFTER ACTION.  
Outer clear zone.  
Inner granular zone  
(discharge of zymogen).

FIG. 75.—SEROUS GLAND. PAROTID; TEASED PREPARATION. (Langley.)

the lobule gradually clears up, the nuclei and borders of individual cells become distinguishable, the granular zone becomes narrower as the homogeneous zone becomes broader, as if a tide of granular material were set up towards and into the lumen of the lobule.

Similar, if somewhat less distinct, changes have been observed on other serous glands. Alcohol-carmine preparations of the rabbit's parotid show that in the gland which has been caused to secrete by excitation of the sympathetic, the alveoli and individual cells are smaller and more stainable, and the cell-nuclei, which in the resting gland are more scattered and irregular in outline, are now of circular outline, exhibiting a clearer structure with more distinct nucleoli. Heidenhain has moreover made out on the parotid of the dog that similar changes of structure are effected by stimulation of the sympathetic,

although no actual secretion is discharged from a cannula in the duct.

Observations on the parotid gland during actual secretion are very difficult; the best that can be done is to examine teased preparations just excised from living glands at rest or after action. This has been done by Langley, who, from a succession of preparations, has made out that clearing of the alveoli and discharge of granules takes place in this case as in that of the pancreas. It may be remarked as a difference of detail, however, that during rest no distinction of zones exists as in the case of the pancreas, but that the whole area of a lobule is granular; zones—viz. an outer clear and inner granular—appear only after secretion has made some progress.

Turning to mucous glands, we find essentially similar changes taking place as the consequence of secretory activity, with certain



Stainable demilunes (= *young protoplasmic cells*).  
Unstainable mucous cells (= *adult cells*).

Increase of demilune cells.  
Destruction of mucous cells.

FIG. 76.—MUCOUS GLAND. DOG'S SUPRA-ORBITAL; ALCOHOL-CARMINE PREPARATION.  
(Heidenhain and Lavdovsky.)

differences of detail. These changes are best studied in stained sections of alcohol-hardened glands, *e.g.* of the supra-orbital and submaxillary glands of the dog. In the alveoli of a resting gland so prepared, two kinds of cells are seen, viz. full-grown large mucous cells, and crescentic clumps of as yet undeveloped cells. The large mucous cells are distended with mucin, which does not stain, and which forms a broad internal zone nearest to the lumen; the outer portion of these large cells nearest to the basement membrane is the stainable protoplasmic zone analogous with that which we have just recognised in the case of serous glands; the crescentic masses are also composed of protoplasm, and stain even better than the protoplasm of the full-grown cells. The appearance of an alveolus of a gland which has been in active and prolonged secretion presents a marked contrast; it is shrunken and as a whole much more deeply stained, so much so that a 'loaded' and a 'discharged' section can be at once



distinguished by the naked eye; the large mucous cells have disappeared, so have the crescentic masses; the cells within the acini are of one uniform kind, stainable, and therefore to be considered as protoplasmic throughout, of moderate size, with central rounded nuclei. In a less extreme case the crescentic masses may be enlarged with increased number of nuclei—an indication of cell multiplication. These appearances are taken to be evidence of (a) destruction of adult mucous cells, together with (b) growth and development of new cells derived from the crescents. According to Langley's recent observations, the distinction between an inner granular and an outer clear zone can be made out on freshly teased mucous gland, though less sharply than on serous gland; the granules (presumably of mucinogen) are highly hygroscopic, and, unless fixed by strong alcohol or by osmic acid vapour, soon swell up and are transformed into the clear mucous material characteristic of the cells as ordinarily seen in sections.

Comparing the process as it is believed to occur in serous and in mucous glands respectively, we find that in the *serous* gland there is no obvious destruction of particular cells, but a transformation of material in three stages:

1. The formation of protoplasm from blood—*outer zone*.
2. The formation of protoplasm-product (zymogen) from protoplasm—*inner zone*.
3. The formation of ferment from zymogen.

In this transmigration of matter the basement membrane is the receiving surface, the free surface of the secreting cell is the discharging surface.

In the mucous gland we recognise the same three changes. 1. The formation of protoplasm from blood (crescentic clumps of protoplasmic cells); 2. The formation of protoplasm-product from protoplasm (swollen mucous cells); 3. The formation of mucus (in the secretion) from mucinogen (in the secreting cell). But in the case of the mucous cells the transmigration of matter is attended with destruction of adult cells at the discharging surface, with birth and growth of new cells at the receiving surface.

The *secretion of milk* by the mammary gland closely resembles that of saliva by a mucous salivary gland; the mammary discharge is, however, a more irregular process, establishing itself only with parturition, and involving a far more copious secretion of organic matter than any other secretion. A healthy wet nurse yields about 1 litre of milk per diem, containing upwards

of 100 grammes of solids. In correspondence with this great amount, the changes in the gland are proportionately active, the epithelial cells lining the acini breaking down rapidly, and equally rapidly growing up. In the dormant gland of the non-pregnant female the acini are almost blocked by small granular cells; in the active gland the appearances vary according as the gland or portion of gland is full or recently emptied by suckling; in the full state the lining cells are swollen and protruded, and droplets of fat are a prominent feature in the cells, which have a ragged appearance, as if breaking down as well as discharging fluid secretion; in the empty state the lining cells are flattened or more or less cubical cells. In the change from the dormant to the active condition of the gland, the gland-cells that filled the alveoli undergo rapid increase and disintegration, giving rise in

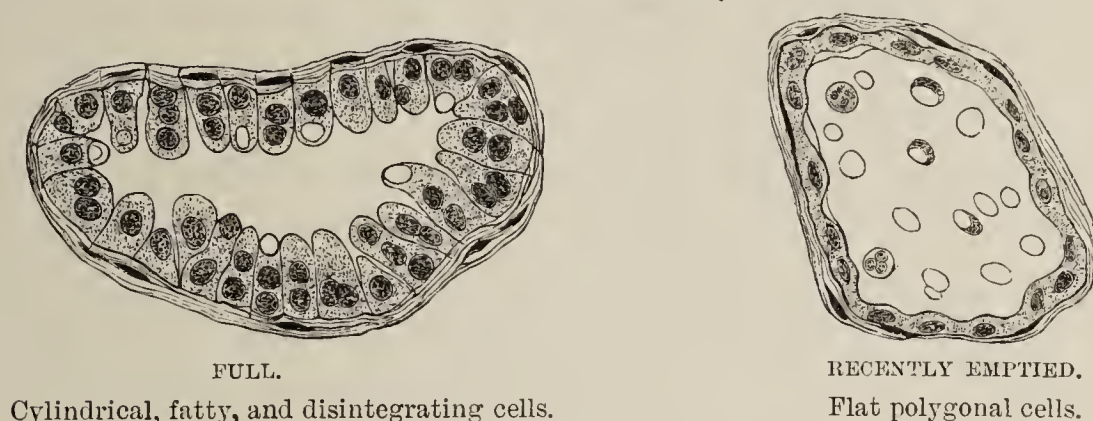


FIG. 77.—MAMMARY GLAND OF BITCH DURING LACTATION. (Heidenhain.)

all probability to the colostrum corpuscles which are a characteristic feature of milk during the first days of lactation. This early milk, or 'colostrum,' contains accordingly a high percentage of proteid, and, unlike ordinary milk, coagulates on boiling.

Pilocarpine, which has so pronounced an effect upon the salivary and sudoral secretions, has little or no influence upon the secretion of milk.

*Other physical changes associated with salivary secretion.*—The act of secretion is attended with—(1) increased blood-supply, (2) expenditure of gland-substance, to which are to be added (3) evolution of heat, (4) increased production of  $\text{CO}_2$ , and (5) electromotive changes.

The evolution of heat has been gauged by comparing the temperature of arterial blood (in the carotid) with that of venous blood, and of saliva, the observations having been taken on the dog's submaxillary. During repose of the gland no constant difference of temperature was found between the various fluids named, but when the gland was in action the temperature of venous



blood and of saliva exceeded (by as much as  $1.5^{\circ}$ ) that of arterial blood, *i.e.* the active gland is a source of heat (Ludwig, 1857).

We have seen that the percentage of  $\text{CO}_2$  is smaller in the venous blood of active than in that of resting gland; an increased production of  $\text{CO}_2$  in an active gland is only to be recognised by taking into account the great amount of blood flowing through an active gland, *i.e.* about fourfold that flowing through a resting gland. Calculations so made of the absolute amount of  $\text{CO}_2$  produced show that it is greater during activity than during repose, although the percentage of  $\text{CO}_2$  in the blood is lower.

**Influence of the nervous system.**—The effect of nerves upon salivary secretion is of great importance, inasmuch as it has furnished the foundation of all definite knowledge relating to the nervous mechanism of secretion.

The chief facts have been ascertained by experiments on the submaxillary gland of the dog, and are confirmed by experiments on the same gland of other animals, as well as upon the parotid.

The submaxillary gland of the dog is innervated through two channels, viz. the chorda tympani and the cervical sympathetic. We have already learned that, as regards action upon the vessels, the chorda tympani dilates, and the sympathetic constricts them, the nerves thereby exercising an indirect influence upon the secreting cells. We have now to learn that both these

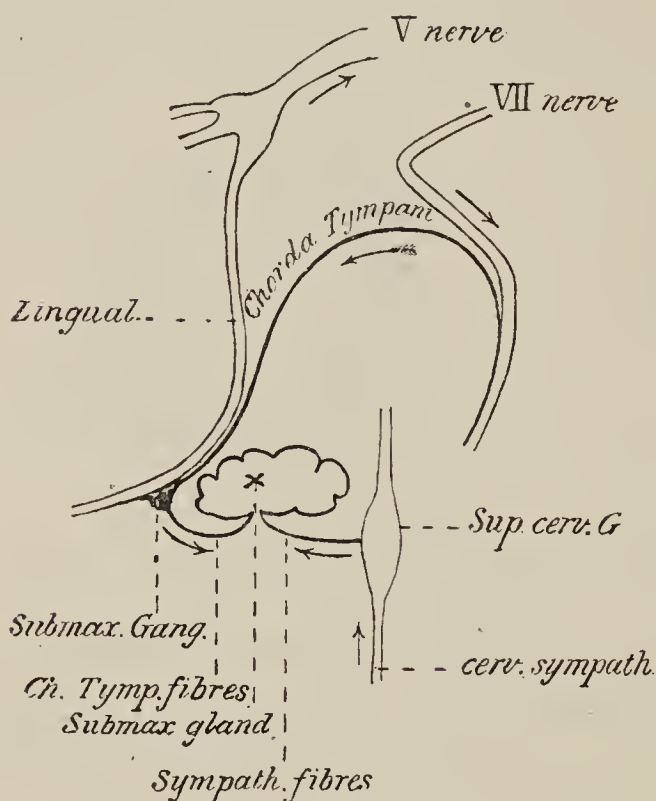


FIG. 78.

Diagram to illustrate the nervous channels to the submaxillary gland of the right side. Chorda tympani fibres pass to it through the submaxillary ganglion. Sympathetic fibres reach it by branches from the superior cervical ganglion which accompany the arteries of the gland. The 'recurrent' chorda fibres alluded to on p. 183 run down the lingual nerve and up again, forming loops.

nerves also act directly upon the secreting cell. Proofs of this:—

1. In the first place, we have to recognise that secretion is the act of the secreting cell, and not the passive filtration of fluid through its substance under the influence of blood-pressure. Ludwig showed, by means of manometers inserted in the salivary duct and in the carotid artery, that salivary pressure in

the duct may reach to double the amount of the arterial pressure in the carotid. The greater pressure cannot be derived from the lesser, *i.e.* secretory pressure cannot be derived from arterial pressure, but must take origin from the secreting cell. This fundamental point is confirmed by other, if less cogent, experiments.

2. Immediately after death, even on a head separated from the body, a little salivary secretion can be brought on by exciting the chorda tympani. The observation is, however, less convincing in the fact than in the account of the fact—only a drop or two of saliva can be thus obtained, and room is left to imagine that this minute amount is squeezed out rather than secreted.

The *post-mortem* proof of direct secreto-motor nerves is more satisfactory in the case of the sweat-glands; stimulation of the peripheral end of a kitten's sciatic will cause beads of sweat to appear on the pad for as long as half an hour after death, or even after amputation of the limb.

3. A dog is poisoned by a suitable dose of atropine, *i.e.* one sufficient to arrest secretion; the peripheral end of the chorda tympani is stimulated, the vessels dilate, but no secretion ensues. This experiment proves that secretion is not the necessary physical consequence of increased blood-supply, since the latter can be obtained without the former.

Accepting as proved the direct action upon the secreting cell of secreto-motor nerves, we must still bear in mind that there is normally an association of vaso-dilatation with increased secretion and of vaso-constriction with diminished secretion. There can be no doubt that increased blood-flow increases secretion, and that diminished blood-flow diminishes secretion. Normal variations of function never occur otherwise; for instance, when a horse begins to eat, saliva flows copiously, and the blood-flow through the active gland is fourfold that which takes place through the resting gland. Can the effect of circulation upon secretion be proved? Increased secretion due to increased blood-flow is not, in the case of the submaxillary gland, definitely ascertained, because the means taken to increase the blood-flow (stimulation of the chorda) at the same time directly increases the secretion. But the opposite relation is definitely ascertained, *viz.* diminished secretion due to diminished blood-flow. If during excitation of the chorda tympani, with consequently dilated vessels and copious salivation, the sympathetic be strongly stimulated, the vessels contract and the flow of saliva is arrested; the sympathetic, as we shall see, is secreto-motor, not secreto-



inhibitory; the arrested flow of saliva must therefore be the consequence of arrested blood-flow.

We have no difficulty in admitting the conclusive character of the experiments establishing the secreto-motor action of the chorda tympani; but in consequence of the above-mentioned vascular phenomena, the proof is less clear for the sympathetic. Yet, attentively regarded, the proof is conclusive enough:—

1. We have seen that changes of structure signifying glandular activity are produced by excitation of the sympathetic, even when no flow of secretion manifests itself (p. 176). In other words, the sympathetic does not arrest glandular action, but excites it.

2. By properly graduated stimulation a flow of saliva can be produced, (a) by excitation of the chorda tympani alone, (b) by excitation of the chorda tympani and sympathetic simultaneously. In the first case the saliva is watery, containing very little organic matter; in the second case it is much thicker, and contains a much larger amount of organic matter. In other words, excitation of the sympathetic, so far from counteracting excitation of the chorda, has supplemented it, and to the water has added organic matter.

3. By a variation of the preceding experiment, the possible interference of vaso-constriction is avoided. The chorda tympani is stimulated (a) before, and (b) after, a period of sympathetic stimulation. The saliva secreted during the first period is thin, that secreted during the second period is comparatively thick, *i.e.* the sympathetic excitation has added organic matter to the water.

4. Atropine administered in such dose as to abolish the chorda effect leaves the sympathetic excitability intact, and excitation of the sympathetic now produces a scanty thick secretion.

The general conclusion which has been drawn is that both nerves are secreto-motor, that the chorda tympani contains a predominance of fibres provoking discharge of water, while the sympathetic contains a predominance of fibres provoking the formation and discharge of organic matter; to this last kind of fibres the term 'trophic' has been applied, but they are more correctly speaking katatrophic or katabolic.

To sum up—*The chorda tympani is vaso-dilatator and secreto-motor of water. The sympathetic is vaso-constrictor and secreto-motor of organic matter.*

Similar experiments have led to similar conclusions with reference to the parotid gland, although, owing to anatomical conditions, this gland has been less frequently studied. The sym-

pathetic is the vaso-constrictor and secreto-motor ('trophic') nerve; the nerve of Jacobson is the cerebral or vaso-dilatator and secreto-motor (water) nerve.

*Reflex salivary secretion.*—The effects just described are those obtained by the experimental stimulation of efferent nerves. The natural secretion of saliva is a reflex act involving afferent nerves, nerve-centre, and efferent nerves, the nerve-centre being situated in the spinal bulb, and the afferent channels being the nerves of taste more particularly, but also other nerves, namely, those from the nose, and eye, and ear, and stomach. A reflex flow of saliva may be caused through the nerves of taste by a 'tasty' morsel in the mouth, or through the nerve of common sensation by the mechanical excitation of a bit of india-rubber, or by the irritation of sore gums, or of a bad tooth; it may be caused through the nerves of smell or of sight, by the odour or by the appearance of savoury food; it may be provoked by material in the stomach, or by a gravid uterus; it may be provoked or suspended by any passing thought or emotion; fear dries the mouth; the mere thought of saliva calls forth saliva in abundance.

*The submaxillary ganglion.*—Experiments are on record showing that after double section of the lingual nerve close above and far below the submaxillary ganglion, excitation of the distal end of the bit of nerve attached to the ganglion can give rise to a flow of saliva (Bernard), and this result was offered in proof of reflex action of a peripheral ganglion. There is no doubt about the fact, but the conclusion is questionable; the result is probably due to an excitation of 'recurrent' chorda fibres in the lingual nerve (Schiff).

**Particular kinds of saliva.**—We have seen that there are several kinds of saliva—submaxillary, parotid, sublingual—according to the gland from which it comes; mucous and serous, according as a gland secretes mucin or albumin with the water which all the glands secrete. The separate salivæ are obtained by catheterisation of the several ducts, *i.e.* by inserting a cannula into Stenson's duct for parotid saliva, into Wharton's duct for submaxillary saliva, into the ducts of Rivinus for sublingual saliva.

From an experimental point of view, two different kinds of saliva have been distinguished, more particularly in the case of the dog's submaxillary, according as the secretion has taken place during excitation of the sympathetic or of the chorda tympani. The former, or *sympathetic saliva*, is extremely thick, viscid, and rich in organic matter, and scanty in quantity as compared



with the second, or *chorda saliva*, which is far more abundant and watery—in short, the former is a concentrated, and the latter a dilute secretion. This difference is not, however, of a constant character; thus, from the submaxillary of the dog and of the rabbit sympathetic saliva is thick, chorda saliva is watery, but on the cat both are watery, and the difference, if any, is the other way, sympathetic saliva being usually the more dilute. In relation with nervous action a third kind of saliva remains to be mentioned, viz. *paralytic saliva*, which is extremely watery and copious; it comes on a few hours after section of the chorda tympani, and in the course of a few days the paralysed gland is distinctly smaller and lighter than the gland of the opposite side; at the end of a month or six weeks the weight of the atrophied gland may have fallen to half that of the normal gland. It is to be remarked that these effects are due to section of the cerebral nerve alone, and that they do not ensue upon division of the sympathetic nerve (the so-called ‘trophic’ nerve), nor upon extirpation of the superior cervical ganglion. The histological characters are (according to Langley) those of resting gland.

**Effect of drugs.**—The two drugs that have been most fully studied as regards their effect on salivary secretion are *atropine* and *pilocarpine*. Attention has also been given to the action of physostigmin, of nicotin, and of muscarin. Atropine—as small a dose as 1 milligramme on man—makes the mouth dry. On the dog, according to Heidenhain, 10 to 15 milligrammes arrest the secreto-motor action of the chorda tympani, without interfering with its vaso-dilating action, or with either the vaso-constrictor or the secreto-motor action of the sympathetic, and the dose was increased to as much as 100 mg. without implicating the sympathetic. Atropine is therefore regarded as producing an interruption of function at the junction between chorda nerve-fibre and submaxillary secreting cell. On the cat this difference, as Langley has shown, does not hold good; in this case both sympathetic and chorda saliva are watery, and atropine (at a somewhat higher dose, it is true) abolishes the secretory effect of sympathetic as well as of chorda stimulation.

The paralysing effect of atropine is counteracted by pilocarpine; and, *vice versa*, the exciting effect of pilocarpine is counteracted by atropine. Pilocarpine in very small amount (1 to 5 milligrammes on the cat) causes a copious flow of saliva, but in large doses (100 to 200 mg.) arrests it and paralyses the secretory nerves. The antagonism between atropine and pilocarpine thus holds good

only for small doses. The exciting action of pilocarpine is in part central, in part peripheral; it is partly central, since section of the chorda tympani diminishes the exaggerated secretion; it is partly peripheral, since the secretion continues excessive after both the secreto-motor nerves have been divided. *Muscarine*, *nicotine*, and *physostigmine* are very similar in their action to pilocarpine; in small doses they excite, in large doses they abolish secretion, and, as in the case of pilocarpine, the exaggeration of secretion produced by small doses is suppressed by atropine.

## II. Gastric Juice

**Physiological anatomy of the stomach.**—The stomach is a secreting churn—secreting by virtue of the glands of its mucous membrane, a churn by virtue of its layers of muscle.

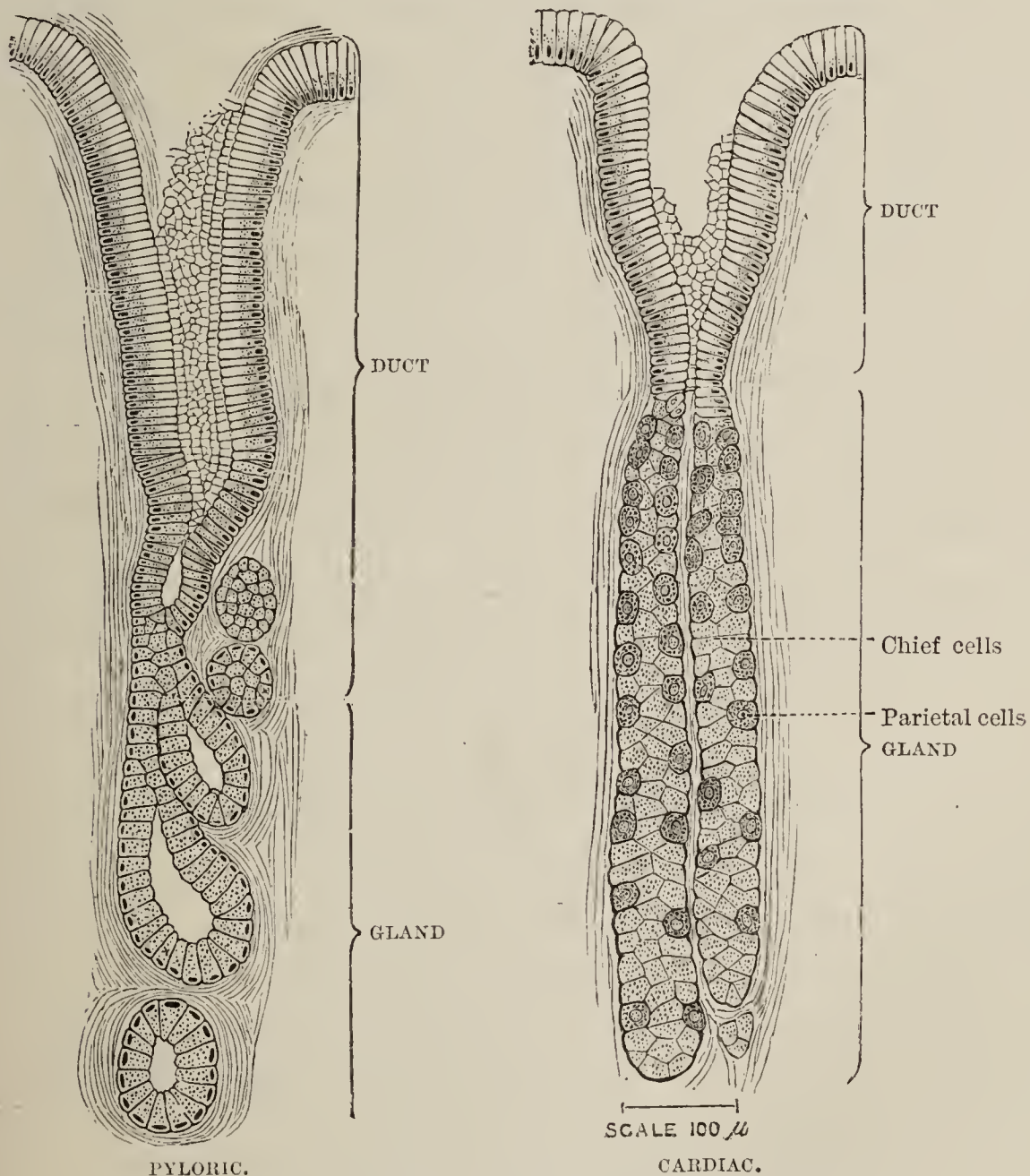


FIG. 79.—GASTRIC GLANDS. (From Heidenhain after Ebstein.)

The mucous membrane of the stomach differs in structure in the cardiac and pyloric portions of the viscus. The glands of



the cardiac end are comparatively long-branched tubules opening on the surface by a short neck or duct; those of the pyloric end are comparatively short-branched tubules opening on the surface by a long neck or duct. In both kinds of gland the neck is lined with columnar epithelium like that lining the free surface of the mucosa; the gland proper is in the pyloric region lined by granular cubical cells, in the cardiac region by similar granular cubical cells (chief cells), and by ovoid parietal cells (accessory cells). We shall see grounds for admitting that these two kinds of cells contribute different constituents to the gastric secretion, the chief cells being the source of pepsin, the accessory cells being the source of acid.

**Gastric juice.**—The secretion of the gastric glands is a colourless acid fluid of low specific gravity (1,002), containing *pepsin* and other ferments, and possessing a slight antiseptic action. The acidity of gastric juice is due to free hydrochloric acid, amounting to 2 per 1,000 of the fluid. The total daily secretion of fluid has been estimated at  $\frac{1}{10}$  the body-weight, or 7 litres; it must not, however, be supposed that this is discharged from the body; it is reabsorbed in the small intestine. The principal ferment, pepsin, acting in concert with hydrochloric acid, converts proteids into peptones. Another ferment, the rennet ferment, brings about the curdling of milk.

*Nature of the gastric acid.*—It was formerly a much-debated question whether hydrochloric or lactic acid is the normal acid of gastric juice; now, however, it is accepted to be the former, for these among other reasons: 1. Separate estimates (*a*) of the chlorine, (*b*) of the bases present in gastric juice, show that the amount of chlorine present is in excess of the amount that could be combined with the amount of basic elements; such excess can be present only in one form, viz. as hydrochloric acid. 2. Several of the aniline dyes, such as methyl violet, are altered in colour far more readily by mineral than by organic acids; gastric juice is found to act like a mineral acid in this respect. 3. A mineral acid (*e.g.* hydrochloric acid) is very soluble in water, hardly at all soluble in ether; an organic acid (*e.g.* lactic acid) is soluble in water and in ether. If fresh gastric juice is shaken up with ether, the latter is subsequently found to contain hardly any acid, from which the conclusion is drawn that the acid in gastric juice is not an organic, but a mineral acid. 4. Uffelmann's test for lactic acid gives a negative result with fresh gastric juice. The test-fluid is changed from blue to yellow by lactic acid,

but simply decolorised by hydrochloric and other mineral acids.<sup>1</sup>

**Action of gastric juice.**—The action of gastric juice has been studied by observing the changes that swallowed food undergoes in the stomach, or by following the transformation of food-stuffs that is effected by gastric juice out of the body. For the latter purpose we may use natural gastric juice obtained from a fistula, or artificial gastric juice made by extracting the mucous membrane of a recently killed animal with dilute hydrochloric acid (1 per 1,000) or with glycerine. The action of gastric juice bears exclusively on proteids, which are dissolved and converted into peptones. If we examined (*a*) the contents of the stomach during the digestion of proteid, (*b*) the blood of the portal vein, we should find in the portal blood no trace of peptone, in the contents of the stomach little or none. This apparent paradox depends on these two facts: 1. That peptone is absorbed as fast as it is formed; 2. That it is reconverted into serum-albumin as fast as it is absorbed. Otherwise expressed, albumin traverses the mucous membrane temporarily disguised as peptone; it is albumin before passage, albumin after passage, peptone only during passage.

All descriptions of the successive changes that a proteid undergoes when acted upon by pepsin are based upon prolonged artificial digestions made '*in vitro*.' Without entering upon a critical account and comparison with each other of the different views of the process which have been advanced, we shall in this place confine our attention to the easily verifiable results of one representative experiment. A handful of boiled fibrin is allowed to digest in a warm chamber at 35° to 40° in a litre of dilute hydrochloric acid, strength 1·5 per 1,000, to which has been added a small quantity of 'gastric glycerine' (*i.e.* a glycerine extract of the mucous membrane of dog's or pig's stomach). We have thus all the conditions necessary to the accomplishment of an artificial digestion, viz. fibrin as the proteid to be digested, an optimum temperature, a dilute acid, and ferment (pepsin) in the gastric glycerine. A few hours later, less or more, in proportion with the power of the glycerine extract employed, we shall by appropriate tests find in solution at least three forms of proteid. (1.) By carefully neutralising a portion of the fluid with sodium carbonate, a precipitate will be obtained. This neutralisation

<sup>1</sup> Uffelmann's fluid.—3 drops of liq. ferri perchlor. (B.P.) in 100 c.c. carbolic acid, 1 per cent. solution.



precipitate is *acid albumin*, otherwise called *syntonin*. (2.) On adding strong  $\text{HNO}_3$  to another portion of the fluid, a precipitate is obtained, which disappears on heating and returns on cooling. This precipitate is *albumose*, sometimes called *propeptone*, or as a more general term, *proteose*. (3.) A third portion of the fluid is boiled with excess of ammonium sulphate (by which all proteids except peptone and a trace of albumose are precipitated). The fluid is filtered, and to it are added a trace of copper sulphate and an excess of sodium hydrate; a rose-colour appears, indicating the presence of *peptone*.

We have learned from this experiment, firstly, the conditions necessary to artificial digestion; secondly, the changes that a solid proteid undergoes in the process. If any one of the conditions be unfulfilled, digestion does not take place; if the experiment be repeated, omitting the acid, or omitting the ferment, or keeping the mixture cold without omitting any ingredient, digestion does not take place. Or if it be repeated with ferment that has been boiled, the same negative result occurs, this, indeed, being the fundamental test informing us that a ferment is concerned in the change.

As regards the phases of the transformation, we learn that proteid is first dissolved, becoming acid-albumin or syntonin, and that, passing through a preliminary stage (propeptone or albumose), it finally becomes peptone. The sequence reminds us of the starch-to-sugar change effected by ptyalin; dextrin is the name of the intermediate station between starch and sugar, albumose is the name of the intermediate station between proteid and peptone; and as there are grounds for admitting that the intermediate between starch and sugar is a series of dextrans, rather than one definite dextrin, so there are reasons for presuming that the intermediate between proteid and peptone is a series of albumoses, rather than one definite albumose. The analogy is drawn still closer when we recognise that the final product of a gastric digest is composed of albumose and peptone, as the final product of a salivary digest is composed of dextrin and sugar.

Some of the obscurity surrounding the subject of proteid digestion is due to the indefinite use of the term 'peptone.' Formerly it included substances which are now called 'albumose'; old-style peptone is a mixture of albumose and peptone, which when in solution can be separated by saturation with ammonium sulphate, albumose being precipitated thereby, while peptone remains in solution. Peptone, as the term is now used, is restricted to that substance or substances

which remain in solution after treatment with ammonium sulphate.<sup>1</sup> Commercial peptone contains a large quantity of albumose, but very little true 'peptone.'

*Albumoses.*—We may take this opportunity of emphasising the great importance of these intermediate digestion-proteids—collectively termed albumoses—by a brief enumeration of facts.

The most rapidly fatal of animal venoms, cobra poison, owes its property to the presence of an albumose. Substances produced by the proteolytic action of pathogenic microbes, and which have recently been found to promote immunity from the further action of such microbes (Wooldridge, Hankin), belong to the family of albumoses. And we may also refer back to the fact, already stated, that albumoses injected into the vascular system produce profound changes—loss of coagulability, fall of blood-pressure, and death. Enough has been said to show that proteids are also protean, and that they may offer many kinds of opportunities 'for different kinds of chemical intercourse.' One kind of proteid may be nutrient, another may be poisonous, another may be protective from further infection; the same proteid (albumose) that nourishes the body if it enters and is modified by the intestinal epithelium, fails to nourish a tissue with which it comes into immediate contact (see p. 99), and proves actually poisonous when directly injected into the blood-vessels. For reasons of this order, and on account of the great importance which the further study of albumose is assuming, we append a summary account of the principal albumoses, and of the procedure by which they may be separated from each other, although it should be stated that their distinguishing properties are not such as to entitle them to be recognised as distinct bodies in the eyes of a chemist.<sup>2</sup>

*Pepsin.*—We have learned that a ferment—pepsin—is the essential constituent of gastric juice. What are the source, properties, and methods of estimation of this important agent? As regards its *source*, it is formed by the chief cells of the cardiac and of the pyloric glands. The nature of the evidence upon which this statement rests is as follows:—The cardiac glands contain two kinds of cells—the chief cells and the accessory cells; the pyloric glands contain only one kind of cells—the chief cells. The secretion of the cardiac end of the stomach contains pepsin and acid; the secretion of the pyloric end of the stomach contains only pepsin. Combining these data, it follows that the chief cells contribute pepsin, while the accessory cells contribute acid to the complete secretion.

<sup>1</sup> But see Appendix, p. 599, footnote.

<sup>2</sup> Appendix, p. 600. The synopsis of the principal proteids may with advantage be consulted at this stage.



In the account given below of the alterations of structure coincident with secretory activity, it will be stated that in the cells a granular substance is present which is not pepsin but *pepsinogen*. The evidence upon which this statement rests is the following : — A fresh mucous membrane is extracted with glycerine, and the extract is found to have a very slight activity ; another portion of the same mucous membrane, previously submitted to the action of HCl (.15 per cent.), yields a glycerine extract of considerable activity. The difference is attributable to a formation of pepsin from pepsinogen under the influence of weak acid.

*Estimation of pepsin.*—Pepsin, like other ferments, has never been completely isolated, but has only been obtained in solution. Like other ferments, it has indefinite power ; presumably minute quantities effect the transformation of a large bulk of proteid, provided that the resultant peptone is not suffered to accumulate ; a single dog's stomach, treated with 200 litres of acidulated water digested 75 kilos of albumen in fifteen days. The quantity of pepsin in a mucous membrane cannot be directly measured, but only indirectly estimated by comparing with some arbitrary standard the activity of a given weight of mucous membrane. For this purpose *Grützner's method* is the most serviceable. It is as follows:—Boiled fibrin is stained with an ammoniacal solution of carmine (.2 per cent.), and well washed first with water, then with dilute HCl (.1 per cent.), until the washings are completely colourless. A portion of this fibrin acted upon by a gastric extract will be dissolved, the carmine belonging to the dissolved fibrin will be set free in the fluid, and tinge it more or less deeply in proportion with the amount of fibrin dissolved. The depth of tint is estimated by comparing it with that of a series of ten test-tubes, I to X, containing :

I	II	III	IV	V	VI	VII	VIII	IX	X	
19.9	19.8	19.7	19.6	19.5	19.4	19.3	19.2	19.1	19.0	.....c.c. water.
.1	.2	.3	.4	.5	.6	.7	.8	.9	1.0	.....c.c. .1% carmine-glycerine,

forming thus a colour-scale. If, for instance, we have to compare the digestive activity of two fluids, we may leave, say, 20 c.c. of each, acidified with HCl to the proper degree, in a warm chamber with 1 c.c. of carmine fibrin. If at the end of a given period the tints of the two solutions are judged to correspond with III. and with VI. of the colour-scale, we conclude that the second solution is twice as active as the first, and presume that it contains twice as much pepsin. Judgment must, of course, be exercised in the use of the method, and the conditions of

observation so adjusted that convenient tints within the scale may be produced in a given time.

The relative amounts of free pepsin and of fixed pepsinogen present in a mucous membrane cannot be precisely determined, but only roughly estimated by comparing the digestive power of simple glycerine extract with that of the subsequent acid extract; and to obtain comparable values it is necessary to adopt regular periods of extraction; *e.g.* to admit for the pepsin extract the product of eight days' extraction of a given weight of dried and pulverised mucosa with 100 times its weight of glycerine—for the pepsinogen proportion the product of one day's extraction of the residue with 1,000 times its weight of dilute HCl (·15 per cent.). The relative activities of the two extracts can then be compared by Grützner's method as just described.

Langley estimates the relative amounts of pepsin and pepsinogen present in a gastric extract by comparing its activity before and after it has been submitted to the action of an equal bulk of  $\text{Na}_2\text{CO}_3$  (1 per cent.) for 15 to 30 seconds. If, for instance, the peptic activity of an extract is equal to  $x$  before  $\text{Na}_2\text{CO}_3$ , and to  $v$  after  $\text{Na}_2\text{CO}_3$ , he concludes that the solution contains equal amounts of pepsin and pepsinogen. If it is  $x$  before  $\text{Na}_2\text{CO}_3$  and  $11$  after  $\text{Na}_2\text{CO}_3$ , the proportion of pepsin to pepsinogen would be  $\frac{8}{2}$ , or 4 to 1. It is to be observed that the division made on either of these two methods is a conventional one; in the first method we agree to call what is in the glycerine extract 'pepsin,' what is in the subsequent acid extract 'pepsinogen'; in the second method we agree to attribute the loss of digestive power in consequence of the action of an equal volume of  $\text{Na}_2\text{CO}_3$  during 30 seconds to 'pepsin,' and the digestive power remaining after such action to 'pepsinogen.'

*Other ferments.*—It has long been known that the fourth stomach of the calf causes a rapid coagulation of milk; under the name of rennet an extract of calf's stomach is commonly employed to make 'curds and whey' in the process of cheese manufacture. Hammarsten has shown that, in addition to pepsin, gastric juice (especially that of sucking animals) contains a distinct rennet ferment (rennin or pexin). It has the property of curdling milk with alkaline reaction, like other ferments it is destroyed by boiling, and its essential effect is the production of a clotted body, *casein*, from a precursor existing in solution of milk, '*caseinogen*'; the production of fibrin from fibrinogen by ferment action has already been alluded to as a similar event



(p. 18), and we shall meet with a third similar instance in the case of myosin (p. 320), and in each case the presence of calcium is a necessary condition. Alkalinity during the process of curdling is the characteristic in the rennet action; acid can of itself curdle milk, no ferment action being involved, so that curdling in an acid medium is not demonstrative of the presence of a ferment.

Evidence of the presence of a 'lactic acid ferment' (producing lactic acid from lactose), of a 'glucose ferment' (converting saccharose to glucose), of a 'fat-splitting ferment' (producing fatty acid from fat), has, moreover, been obtained by various observers. But the effects of these are hardly deserving of mention, as they are not characteristic of gastric juice nor significant of its normal action. It is of more practical importance to recognise that gastric juice is antiseptic; left to itself it

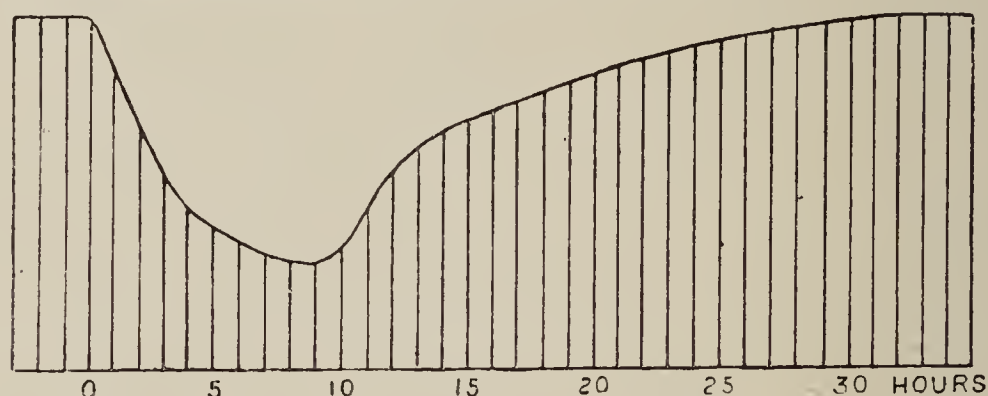


FIG. 80.

Peptic activity of gastric mucous membrane (cardiac) after the ingestion of food. (Grützner.)

resists putrefaction for an indefinite period, and normally it no doubt has a 'sterilising' action upon swallowed materials.

**Changes consequent upon secretory activity**, which are a prominent and well-known event in the salivary glands, have also been studied in the gastric glands, and very similar though not identical changes have been observed. Heidenhain describes the changes occurring in the dog's stomach after food as follows:— During the first six hours the chief cells are enlarged, clouded, and more stainable, the ovoid cells are somewhat swollen. From the 6th to the 9th hour the chief cells shrink more and more, and become more cloudy, the ovoid cells are still swollen. This state lasts till the 13th to 15th hour. From the 15th to 20th hour the chief cells gradually enlarge again and clear up, the ovoid cells shrink, the gland thus resuming its normal appearance during fasting. This account applies to the cardiac end of the stomach; in the glands of the pyloric end, containing only

chief cells, the changes occur somewhat later ; it should be added that the times given above are liable to great fluctuation.

Comparing the appearances of the chief cells with the peptic power of the mucous membrane at various periods after a meal, it is found that with the large, clear, relatively unstainable cell, the digestive power of a given weight of mucous membrane is three times as great as when the chief cells are in the shrunken, clouded, and more stainable state. Heidenhain deduces from these observations conclusions essentially identical with those previously given in the case of salivary glands. During the fasting state the zymogen (in this case pepsinogen) is formed from protoplasm and stored in the chief cells. During secretion three processes are simultaneously progressing :—(1) The new growth of protoplasm. (2) Continued production of pepsinogen. (3) Discharge of pepsinogen as pepsin.

Heidenhain's observations were made on stained sections of alcohol-hardened stomachs of dogs. Langley has made similar observations on the fresh stomachs of frogs and of mammalia, and comes to a similar conclusion. In the fasting state (18 hours after a meal) he finds the glands uniformly granular throughout ; as digestion proceeds he observes the formation of zones—outer clear and inner granular—and the progress of a tide of granules towards the lumen of the gland ; he considers that the granules are the actual pepsinogen. Both the above accounts thus attribute the alterations entirely to the outgoing secretion, and not to any incoming absorption of food.

**Influence of the nervous system.**—There is no direct proof of the existence of any regulation of gastric secretion by secretomotor nerves ; neither section nor excitation of the sympathetic or of the pneumogastric nerves produces any alteration of gastric secretion ; yet certain facts suggest the possibility that nerves may take part in the regulation of secretion. The flow of gastric juice is not continuous, but occurs in response to the ingestion of food. No doubt the direct mechanical action of swallowed food may excite secretion from the mucous membrane, but this is not all : secretion has been observed to occur when, owing to stricture of the gullet, no food could reach the stomach. Mechanical excitation of the mucous membrane has been seen to produce secretion of gastric juice—this is a direct action, independent of nerves ; chemical excitation likewise exercises a direct action on the mucous membrane. A perfectly indigestible body, such as glass or india-rubber, by its mechanical



action produces secretion; but a digestible body, such as a bit of meat, produces a far more active secretion, and Schiff has shown that dextrin and meat-extract are particularly active in causing a true peptic secretion from a previously exhausted stomach; alkalies also are powerful excitants of the secretion, and one of the services of saliva is thus to promote the flow of gastric juice. But these various facts contain no *proof* of an action of the nervous system; the effects can be referred to a chemical action of substances absorbed into and carried by the blood, when they are not attributable to direct mechanical or chemical excitation.

A *gastric fistula* is a permanent abnormal opening leading through the abdominal wall into the stomach. Such an opening may be effected by accident or by surgical operation, and can

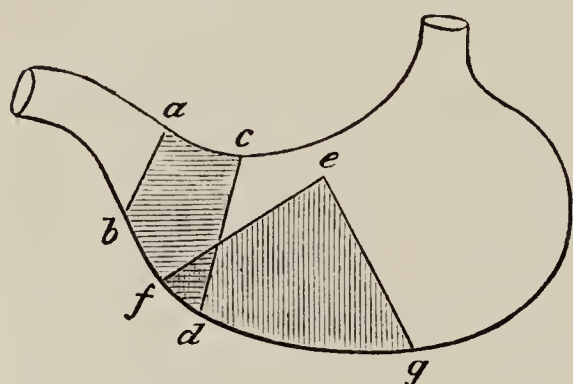


FIG. 81.

Parts of the stomach from which gastric cæca have been formed; pyloric cæcum from *a, b, c, d*; cardiac cæcum from *e, f, g*. (Heidenhain.)

then be utilised for the study of gastric digestion. A *gastric cæcum* is a blind pocket formed by operation and composed of a portion of the stomach. The principle of the operation is as follows. A portion of the stomach, *a, b, c, d*, is included between two incisions, *a b* and *c d*.

The two edges *a b* and *c d* are sewn together and soon heal, so that the continuity of the viscus is restored. One of the two edges of the shaded portion, *a b c d*, which is to form the cæcum, is sewn up and allowed to heal so as to form the bottom of the pocket. The other edge is also sewn up with the exception of a small orifice, the borders of which are fixed to the borders of a corresponding orifice left in the abdominal wall. The final result of the operation is thus an isolated pocket formed of pyloric or of cardiac stomach-wall, with an external orifice through which the mucous membrane can be observed, or materials introduced, or secretions removed. A pyloric cæcum so formed secretes a thick mucous alkaline fluid, which contains pepsin; a cardiac cæcum secretes a thin highly acid fluid, which also contains pepsin. Swallowed food provokes secretion from such a cardiac cæcum, although it does not come in direct contact with it; the secretion thus excited begins in about 15 to 30 minutes, and is considered not to be a reflex effect, as it does not occur unless material is digested

and absorbed by the main stomach ; it seems therefore to be due to a chemical stimulation by blood that is charged with products of digestion.

*Self-digestion post-mortem.*—Under certain circumstances the stomach can digest itself, and the question has been asked, Why does it not normally do so during life? The excised stomach of a recently-fed animal, left in a warm chamber in a sufficient quantity of dilute hydrochloric acid, will become completely dissolved. An animal killed during digestion and examined a few hours later may be found with the stomach perforated by self-digestion, and with more or less extensive destruction of surrounding tissues. The non-digestion of the living stomach by its own gastric juice during life is generally referred to the alkalinity of the blood, but it is a sufficient answer to simply paraphrase the question, and to say that living tissue resists digestive action so long as it is living.

An experiment by Bernard, which is quoted to show that living tissue can be digested, shows no such thing. The experiment consists in the introduction of the legs of a living frog into a dog's stomach through a fistula ; the legs become digested, but we have good reason to say that they are first killed, then digested. Nor is the protective influence of alkaline blood a good and sufficient answer, for it does not apply to pancreatic fluid which acts on proteids in an alkaline medium, yet does not digest the living proteid with which it may come in contact.

### III. *Bile and Pancreatic Juice*

**The liver.**—The chief functions of the liver are (1) the secretion of bile, (2) the formation of sugar. As regards its first function, the liver is an ordinary secreting gland, with a duct opening into the duodenum. As regards its second function, it is a so-called ductless gland, from which elaborated products are carried off by the outgoing venous blood ; and it is a noteworthy point in this respect that the liver is traversed by the main current of incoming material carried by the portal blood. These are the two chief functions of the liver, to which are to be added two minor, or, at any rate, less obvious functions, viz. (3) the destruction of red corpuscles, (4) the formation of urea and of other urinary substances.

We shall at this stage consider only that function whereby a digestive fluid is secreted, namely, the bile ; the sugar function,



which is subsequent to the absorption of food, will be considered under the heading 'Glycogenesis' (p. 213); the urea function, which belongs to elimination of used-up or waste proteid, will be considered in connection with excretion (p. 241).

*Physiological anatomy.*—Three channels enter and leave the liver at the transverse fissure or hilum. The largest of these, the *portal vein*, conveys venous blood from the digestive viscera; material upon which the liver exercises its elaborative action; the smallest, the *hepatic artery*, conveys arterial blood requisite to

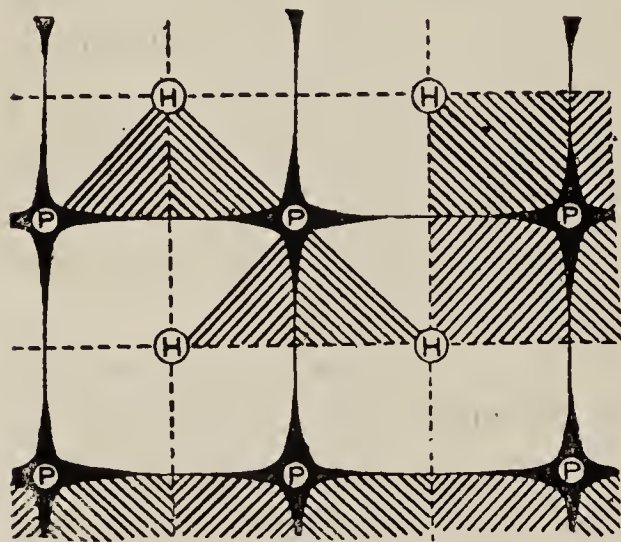


FIG. 82.

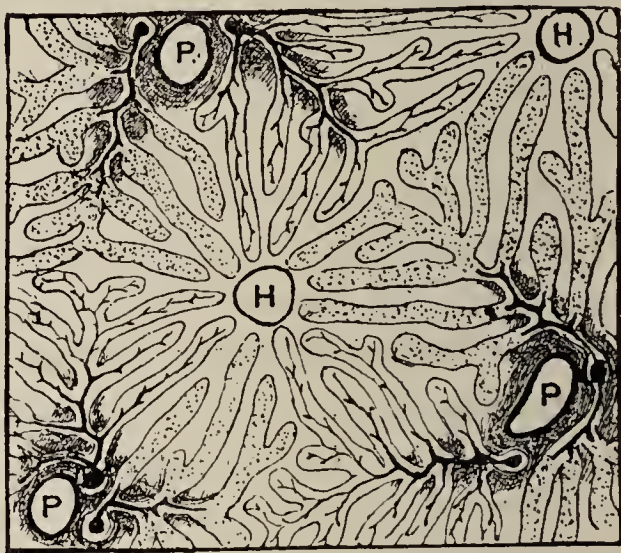


FIG. 83.

Diagrams to illustrate the physiological anatomy of the liver as a biliary gland, according to the investigations of Delépine. In the geometrical diagram (fig. 82) the squares P P P P represent vascular lobules, and the triangles P H H true biliary lobules; H = hepatic veins; P = portal veins surrounded by Glisson's capsule. The striation (indicated only in portions of vascular lobules) denotes the general disposition of the columns of gland-cells, diverging from the dotted lines between H H, and converging to the interlobular lines, P P; the triangle P H H is a true biliary lobule, the triangle H P P is not, but composed of portions of two contiguous lobules, and several such portions enter into the composition of a vascular lobule, P P P P. Fig. 83 is a semi-diagrammatic, to exhibit the tubular structure of the gland and its relation to the bile-ducts; one entire vascular lobule, P P P, and two entire biliary lobules, P H H, are shown, and portions of other lobules; the triangle H P P is seen to be composed of portions of two distinct lobules; an imaginary line joining H H is a line of hepatic divergence, and lines joining P P are lines of portal convergence. Biliary anastomoses are omitted for the sake of distinctness.

the nutrition of the liver substance; the current in these two channels is towards the liver, they penetrate into and ramify in its substance and ultimately form a system of small vessels between adjacent lobules, hence called *interlobular vessels*. The third channel is the *bile-duct*, in which the current of secretion is from the liver, and it is noteworthy that in most animals a *gall-bladder* is present, serving as a storage chamber in which bile, secreted in the intervals of digestion, is kept until required. A fourth system of vessels takes origin within the lobules, from

venules, that are therefore called *intralobular*, and that conjoin to form the *sublobular* and the hepatic veins; the latter emerge from the posterior border of the liver, and carry off blood upon which the organ has exercised its functions.

It is at first sight most natural to regard each lobule as a miniature liver surrounding the intralobular vein and surrounded by interlobular veins; but upon this view the glandular plan of structure is almost entirely lost sight of. In the livers of the lower vertebrates (*e.g.* frog) the tubular type is very clearly exhibited: as seen in section the hepatic gland-cells form double rows, belonging to a network of anastomosing tubules. In the livers of the higher vertebrates, the *obvious* lobulation produced by the disposition of the blood-vessels masks what is probably the *true* lobulation subordinate to the disposition of the bile-ducts; a true biliary lobule is composed of adjacent portions of contiguous vascular lobules; the centre of the latter is a hepatic venule, the centre of the former is a bile-duct, and the gland is really, although not obviously, a mass of anastomosing tubules—the bile-capillaries bounded by liver-cells. These capillaries take origin from the cells themselves by intracellular radicles, and it is to be observed as a noteworthy point concerning the relation of a liver-cell to the surrounding capillaries, that blood-capillaries and bile-capillaries do not lie side by side against the cell, but are as far as possible separate from each other; the bile-channels are between the opposed surfaces of two contiguous cells, the blood-capillaries occupy somewhat larger and longer channels between the adjacent edges of columns of cells.

Physiologically regarded, each liver-cell is the functional centre; hepatic arterial blood feeds it, crude portal blood comes to it, and is modified by it, bile leaves it to do duty in the intestine, hepatic venous blood leaves it to do duty in the tissues of the body. Alterations of structure consequent upon the absorption of food will be considered in connection with ‘glycogenesis.’

**The pancreas**, in its minute structure, very closely resembles a serous salivary gland, being a racemose gland with short tubular acini lined by cubical granular epithelium. It is, as has already been described (p. 175), the chosen gland in which visible tokens of secretory action have been most closely studied. The granular contents of the cells are considered to be the ‘*zymogen*,’ or protoplasm-product which is about to be transformed into ferment, and the experiments which show that the living or perfectly fresh gland contains ferment-forming or zymogenic material, and not



the actual formed ferment, are, in the case of the pancreas, particularly striking and conclusive. The glycerine extract of a perfectly fresh pancreas has hardly any fermentative action; the glycerine extract of a pancreas that has been left for 24 hours in the body of the animal or in a warm chamber is powerfully active. The difference in the two cases is explained by saying that the fresh organ contains zymogen, but not ferment, while in the stale organ the elaboration of ferment from zymogen has taken place. By weak acid, this conversion can be effected at once; a perfectly fresh pancreas, extracted with dilute acetic acid, yields a fluid that has fermentative power.

**Bile.**—*Its composition.* Bile is a reddish-brown or dirty-green fluid of slightly alkaline reaction, with a bitter taste and with a specific gravity of 1,020. The daily secretion of bile is the same as that of urine, viz. about 1,500 c.c. It contains:—

	Water . . . . .	90 to 85 per 100
	Mucous matter . . . . .	
Bile-acids	{ Glycocholic acid . . . . .	
	{ Taurocholic acid . . . . .	
Bile-pigments	{ Bilirubin . . . . .	10 to 15 per 100
	{ Biliverdin . . . . .	
	{ Cholesterin . . . . .	
	{ Lecithin . . . . .	

It does not contain any *albumin*. *Mucus*, or, properly speaking, a mucinoid substance, is generally present in large quantity; it gives to bile a glairy, ropy consistency, and is the cause of the rapid decomposition to which bile is subject, although bile itself possesses antiseptic properties by virtue of other constituents.

The specific biliary substances are the bile-pigments and the bile-acids, the acids being for the most part combined with sodium in the form of salts, as glycocholate and taurocholate of soda, and forming two-thirds of the total biliary solids.

The *bile-acids*, glycocholic and taurocholic, are compounds of glycin and of taurin with cholalic acid. Glycocholic acid boiled with dilute acid or with alkali takes up water and breaks up into glycin and cholalic acid. Taurocholic acid similarly treated breaks up into taurin and cholalic acid. Hippuric acid similarly treated breaks up into glycin and benzoic acid. Glycin is amido-acetic acid, *i.e.* a nitrogenous body; taurin is amido-isethionic acid, *i.e.* a nitrogenous body combined with a sulphur acid. Cholalic acid is a complicated non-nitrogenous body. The

three reactions will be found graphically represented in the Appendix.

*To separate the bile-salts from bile*, the fluid evaporated to one-quarter its bulk is made into a paste with animal charcoal, dried at  $100^{\circ}$ , then pulverised and extracted with alcohol; the bile-salts are precipitable as a crystalline mass by adding ether in excess.

*To separate the two bile-acids from each other* the ether precipitate is redissolved and the solution treated with neutral lead acetate, which gives a precipitate of glycocholate of lead; the precipitate is collected on a filter, and the filtrate, treated with basic lead acetate and ammonia, gives a precipitate of taurocholate of lead, which is likewise collected on a filter. To obtain the acids free of lead the precipitates are respectively dissolved in hot alcohol, and the lead removed by a stream of  $H_2S$ ; from the filtered alcoholic solutions glycocholic and taurocholic acids are respectively precipitated by adding water. Glycocholic acid is best prepared from pig's bile, taurocholic acid from dog's bile.

The *bile-pigments*, as they normally exist in bile, are *bilirubin* and *biliverdin*—biliverdin ( $C_{32}H_{36}N_4O_8$ ) being the more highly oxidised state of bilirubin ( $C_{32}H_{36}N_4O_6$ ). By powerful reducing or oxidising agents less oxidised and more oxidised bodies are obtainable. Nitric acid, as employed, for instance, in the Gmelin test, gives as the final product a yellow substance, *choletelin*, which is the most highly oxidised state of bile-pigment. On the other hand, by the reducing action of sodium amalgam *hydrobilirubin* is produced, the least highly oxidised state of bile-pigment, and identical with the bile-pigment derivatives in urine and in fæces, which are produced by the reducing action of the tissues. These relations are exhibited in the following table:—

$(C_{32}H_{36}N_4O_5 \cdot 2H_2O)$ <i>Hydrobilirubin</i> .		Produced from bilirubin by the reducing action of sodium amalgam. Not distinguished from <i>urobilin</i> and <i>stercobilin</i> and from <i>hæmatoidin</i> .
$C_{32}H_{36}N_4O_6$ )	<i>Bilirubin</i> .	
$(C_{32}H_{36}N_4O_8)$	<i>Biliverdin</i> .	. Produced from bilirubin by oxidation.
$(C_{32}H_{36}N_4O_{12})$	<i>Choletelin</i> .	. Produced from bilirubin by the oxidising action of nitric acid.

*Bilirubin* can be obtained in quantity from 'red' gall-stones. These are pounded, treated with hydrochloric acid to dissolve chalk, and extracted with chloroform; the chloroform solution yields crystals of bilirubin.



*Cholesterin* ( $C_{26}H_{44}O, H_2O$ ) is a monatomic alcohol crystallising in rhombic plates or needles, widely distributed in the animal and vegetable kingdoms, and probably a derivative of proteid, although itself a non-nitrogenous body. The two materials which contain it in greatest abundance are the bile and the brain. White gall-stones in particular, which are a deposit from bile, are almost entirely composed of cholesterin, and the substance is most readily prepared from this source by extraction with hot alcohol. On evaporation, cholesterin is obtained as a crystalline mass which is soluble in alcohol, ether and chloroform; from chloroform it crystallises in characteristic notched plates.

*Tests.*—The presence of *mucus* is recognised by the addition of *acetic acid*, which gives a precipitate.

The presence of *bile-acids* is recognised by *Pettenkofer's test*. This consists in the cautious addition of  $H_2SO_4$ , and of a few drops of cane-sugar solution, or a drop of furfural solution; a purple colour signifies the presence of bile-acids.

The presence of *bile-pigments* is recognised by Gmelin's test. Nitric acid (containing also nitrous acid, as indicated by red fumes) causes a play of colours from green to blue, violet, red and yellow.

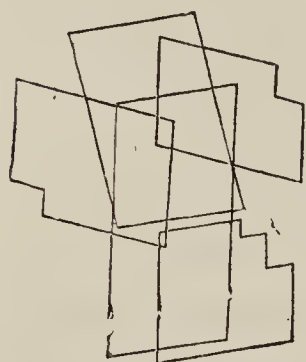


FIG. 84.—CHOLESTERIN.  
NOTCHED PLATES.

*Cholesterin* is recognisable (a) under the microscope by the crystals that are formed in its chloroform solution, (b) by the addition of sulphuric acid to a chloroform solution, giving a bright cherry-red colour.

**Pancreatic juice** is a viscid albuminous secretion of high specific gravity and of alkaline reaction. The amount secreted per diem does not exceed 150 c.c., *i.e.* it is about a tenth of the amount of bile secreted per diem. It contains:—

	per 100
Water . . . . .	90
Albumin . . . . .	} 10
Ferments, namely, lipolytic, proteolytic, and amylolytic . . . . .	
Salts, especially $Na_2CO_3$ . . . . .	

*Tests.*—*Albumin* is recognised by boiling. The amount of albumin is often so large that the fluid is coagulated *en masse*.

*Ferments* are recognised by the changes that take place in appropriate digests, (1) of fat, (2) of proteid, (3) of starch, with pancreatic juice or with pancreatic extract. Fatty acid appears in the first digest, peptone in the second, sugar in the third.

Stale pancreatic juice contains products of the digestion of its own albumin, viz., peptone, leucin, and tyrosin.

#### USES OF THE BILE AND OF PANCREATIC JUICE

Bile is mainly a digestive fluid, *i.e.* it is a secretion to be further utilised in the body, not merely an excretion to be forthwith expelled. The simple anatomical fact that the bile-duct enters the intestine at its very beginning is enough to prove this point. A no less simple physiological experiment confirms the view; if in a dog, for instance, a biliary fistula is made through which bile is diverted, the animal thins rapidly, and by the eagerness he evinces to lick bile from the fistula says plainly enough that bile is wanted; the rapid emaciation in spite of an ample diet, and the fatty nature of the fæces discharged, inform us further that bile is wanted for the *digestion of fat*, since, failing the bile, fat passes undigested and unabsorbed along the intestinal canal. It should be added, however, that the malnutrition caused by loss of bile seems to be far less serious on the human subject; what occurs in a dog does not necessarily apply to man; perhaps the difference may be partly due to the fact that human bile contains very little of the sulphur-holding principle—taurocholic acid—whereas in dog's bile it is *the* bile-acid.

Bile is also to a small extent an excrement. Already during foetal life, when it can have no digestive office, it is discharged into the intestine, where it forms part of the excrement termed *meconium*. In the adult, most of the bile secreted by the liver does its work in, and is re-absorbed by, the intestine; but a small amount is not so re-absorbed, and escapes with the fæces, and another small amount, although re-absorbed, is separated by the kidneys with the urine. These excrementitious substances of bile form, especially the colouring matters of fæces and of urine, *stercobilin* in the former, *urobilin* in the latter, both these substances being indistinguishable from *hydrobilirubin* (= reduced bilirubin).

We are not, however, at this stage concerned with the excretory action of the liver, but only with its secretory action, in so far as bile is a digestive fluid, and shall therefore only allude to the fact that, as an excretory organ, the liver forms substances that find their way into the blood, and are separated from it by the kidneys. We shall see in a future section that this physiological association between kidney and liver as excretory organs



applies to urea and to uric acid, which are probably formed in the liver and carried away by the hepatic blood, as well as to bilirubin, which is carried away by the bile (p. 241).

In addition to its principal action upon fats, the bile assists in digestion by precipitating syntonin, albumose, and pepsin, thus preparing the gastric chyme for the digestive action of pancreatic juice; bile also promotes the digestion of starch; it stimulates peristaltic movements, and it retards the putrefaction of food in the intestine.

The digestion of fats, and of proteids, and of carbohydrates is promoted by bile, the action on fats being, as already stated, the most important of the three; in each case the bile acts in concert with pancreatic juice, the relative action of the two fluids being essentially:—preparation by bile, completion by pancreatic juice. We have an anatomical token of this partnership in the close proximity, or actual conjunction in one channel, of the biliary and pancreatic ducts; physiological evidence of united action is afforded by the parallelism between the biliary and pancreatic secretions, which rise and fall together (fig. 86), pathological evidence by the fact that fatty stools, due to the non-digestion of fat, may result from liver disease alone, or from pancreatic disease alone, showing us that the healthy secretion of both glands is necessary to the digestion of fat.

Experiments outside the body fully bear out these statements.

1. *Experiments on fats.*—Bile and oil shaken together form a fine and permanent emulsion; this finely divided state is obviously favourable to digestion. Pancreatic juice and oil likewise form a good emulsion; pancreatic juice, moreover, effects an actual division of fat into glycerine and fatty acid. This division is in many ways favourable to digestion. The fatty acid, of itself, favours emulsification, and still more so by virtue of the fact that by uniting with the sodium of the bile-salts it forms soap. The soap so formed not only favours emulsion, but it is itself diffusible through animal membranes, and its presence in their pores enables them to be traversed by fat as such. Thus bile and pancreatic juice acting together promote the absorption of fat by favouring its emulsification, its decomposition, and its saponification. The composition of fats and soaps will be found graphically represented in the Appendix.

*Bernard's observations.*—Very striking evidence of the part played by the pancreas in the absorption of fat was furnished by

Bernard's observations on the rabbit. In this animal the pancreatic duct opens into the intestine about 30 c.m. below the opening of the bile-duct; if a rabbit is killed two or three hours after receiving a fatty meal, and its mesentery examined, the lacteal vessels above and below the opening of the pancreatic duct will be found very different in appearance; in the portion of mesentery connected with the intestine above the duct the lacteals contain clear lymph without fat, while below it they contain white chyle with an abundance of fat; that is to say, fat is being absorbed only in the portion of intestine below the duct after the pancreatic juice has taken effect upon it.

2. *Experiments on proteids.*—The addition of bile to a gastric digest causes a precipitate of syntonin and of albumose, and puts an end to the action of pepsin. In the body these effects are probably the necessary antecedent to pancreatic digestion. The partially digested proteid contained in the acid chyme which is gradually pressed through the pylorus requires to be acted upon by bile before it can undergo further digestion by pancreatic juice; in other words, bile arrests peptic digestion in an acid medium, and prepares chyme for pancreatic digestion in an alkaline medium. This interference with peptic digestion, which in the duodenum is the normal event, can occur as an irregular event in the stomach; bile may be driven



FIG. 85.—TYROSIN AND LEUCIN SHEAVES AND LUMPS.

the wrong way from the duodenum into the stomach, where it interferes with digestion and produces vomiting. Pancreatic juice has a powerful digestive action upon proteids (Corvisart, Kühne), which it owes to the presence of the ferment *trypsin*; under its influence proteids are converted into peptones which are indistinguishable from gastric peptones. This digestive action on proteids differs from that of gastric juice in the following particulars:—(a) It occurs only in an *alkaline* medium. For a gastric digest we employ dilute hydrochloric acid .15 per cent.; for a pancreatic digest the best medium to employ is a 1 per cent. solution of *sodium carbonate*. (b) The proteolytic power of trypsin is greater than that of pepsin; the latter does not carry proteid beyond the peptone stage; the former, besides converting proteid to peptone, can break up peptone, giving rise to *leucin* and to *tyrosin*, and at a latter stage to indol, phenol, and other bodies. (c) As subsidiary



points of difference it may be mentioned, that whereas in a gastric digest acid-albumin is an intermediate product, in a pancreatic digest it is alkali-albumin; and that when fibrin is used for such experiment it swells up before solution in the gastric fluid, while in the pancreatic fluid it is gradually corroded and dissolved. Moreover, the great rapidity with which a pancreatic digest enters into putrefactive decomposition is characteristic; in a few hours it has a repulsive smell, and a drop of the fluid examined under the microscope is seen swarming with bacilli.

3. *Experiments on carbohydrates.*—Bile by itself has little or no digestive action upon carbohydrates; at most it has a slight amylolytic action, converting starch into sugar. The principal digestive agent of starch is pancreatic juice, which acts in this respect far more actively than the most active saliva. Soluble or even raw starch left to digest at 40° with pancreatic juice or pancreatic extract, is rapidly converted into sugar of the maltose variety. The presence of bile seems to promote this digestion; if we compare two digests of starch and pancreatic extract, to one of which bile has been added, this last will be found to contain more sugar than the digest made without bile (Martin).

Reviewing the preceding experiments, we see that the statement made at the outset, to the effect that the combined digestive action of bile and of pancreatic juice is essentially preparation by the former, completion by the latter, is fully justified, and applicable to all three classes of food-principles, to proteids, to fats, and to carbohydrates. Bile contains no ferments. Pancreatic juice contains *lipolytic*, *amylolytic*, and *proteolytic* ferments, acting respectively upon fat, upon starch, and upon proteid.

For the sake of completeness, rather than because we have any reason to believe that the action is normally exercised, it may be mentioned that pancreatic juice or extract possesses the rennet property of coagulating milk; we have, however, no ground for supposing that milk can ever escape coagulation in the stomach, either by rennet ferment or by acid, and be coagulated by pancreatic juice.

**Derivation of the bile-substances.**—The specific biliary substances (pigments and acids) are made by the liver, not simply removed by it ready-made from the blood. This is in contrast with what we shall find in the case of the kidney which simply separates from the blood ready-made urea produced by the tissues. What are the facts upon which the above doctrine is

based? Briefly stated, they are as follows:—After removal of the kidneys, urea accumulates in the blood; *i.e.* it continues to be formed in the absence of the kidney; after removal of the liver, the biliary substances do not accumulate in the blood, *i.e.* the presence of the liver is necessary to their formation.

It must, however, be observed that the contrast is not actually as precise as set forth in the above curt statement. On the one hand, we have to admit that some of the specific urinary constituents are formed by the kidney; on the other, that some of the specific biliary constituents may be formed in the tissues. So that it is a nearer approach to the truth to say that a greater part of the bile-substances are made by the liver, a lesser part may be made by the tissues, while a greater part of the urinary substances are formed by the tissues, a lesser part by the kidneys.

Let us consider the case of the bile-pigments. The liver is one of the organs in which red corpuscles are destroyed, and the hæmoglobin derived from them is the source of the bile-pigment formed by the liver. Acting upon this liberated hæmoglobin the liver forms bile-pigment. We have thus accounted for the greater part of the bile-pigments.

Blood effused into the tissues is found to have formed hæmatoidin by simple lapse of time (*e.g.* in old cerebral hæmorrhages); this is undoubtedly a blood-pigment, and it is indistinguishable from the bile-pigment, bilirubin. Here we have a visible instance which leads us to admit that the tissues themselves, acting upon blood-pigment, can form a bile-pigment. This is a reason for admitting that a lesser part of the bile-pigments may be formed by the tissues.

Bile-pigment differs from blood-pigment in this important particular, that the former does not contain iron. We should accordingly expect to find—and in point of fact we do find—free iron salts deposited in the liver tissue; a slice of liver washed and immersed in a solution of potassium ferrocyanide (10 per cent.) for a few minutes, washed again and left in hydrochloric acid (2 per cent.) for a few hours, turns *blue*, indicating the presence of peroxide of iron. This free iron must come from hæmoglobin, which is broken up, partly in the liver, partly in the tissues generally; in the latter case it exists in the plasma as an albuminate of iron, which in order to give the blue reaction must first be oxidised by combustion or by boiling with  $\text{H}_2\text{SO}_4$ , and it is not improbable that some of the liver iron is formed from this source.



Extirpation of the liver yields very little information as to hepatic function, because cold-blooded animals and birds are alone capable of surviving the operation for any length of time. As regards mammalian animals, we must have recourse to pathological observations. Obstruction of the bile-ducts leads to an accumulation of bile-pigments in the blood and in the lymph (the yellow discoloration so produced is called jaundice); whereas abolition of hepatic function does not lead to such accumulation at all, or causes it in much less degree. In the first case we have the obvious change brought about by an active liver with its duct obstructed; in the second we have the doubtful or negative consequence of diminished liver action. The conclusion is as before, that bile-pigment is certainly and in large quantity formed by the liver, and that, failing the liver, it is possibly, and in small quantity, formed in the tissues of the body.

This amounts to saying that bile-pigment is normally formed by the liver; the association between bile-pigments and bile-acids is so constant, that we are justified in believing that what is true of bile-pigments is also true of bile-acids, and that the latter also are normally formed by the liver, not simply removed by it ready made from the blood.

*Biliary pressure.*—The pressure to which the biliary secretion can reach, as measured by a manometer tied in the duct, is low, not more than  $1\frac{1}{2}$  cm. of mercury. This has important practical bearings, as well as some theoretical interest. It explains to us how slight a cause is sufficient to arrest the flow of bile, and consequently give rise to its re-absorption and to jaundice. A gall-stone in the duct will obviously enough produce the effect, but it may be due to apparently far slighter causes; a swollen and congested state of the mucous membrane of the duodenum is a sufficient impediment, and a frequent cause of jaundice. The re-absorption occurs through lymphatics; in dogs the re-absorbed biliary substances can be detected in the lymph, and they are not found in the blood if the thoracic duct is ligatured.

The theoretical interest attaching to biliary pressure lies in its comparison with blood-pressure. It has been remarked that the maximum biliary pressure of  $1\frac{1}{2}$  cm. Hg is much below ordinary blood-pressure, so that the proof of secretory independence given for salivary secretion does not cover the case of biliary secretion; but it is to be remembered that the hepatic blood-pressure is not that of the arterial system, but only that of the portal vein, which is equal to less than 1 cm. Hg.

There is no experimental evidence of any direct action of nerves upon biliary *secretion*, although the *expulsion* of bile from the gall-bladder into the small intestine occurs in response to stimulation of the duodenum by the acid chyme; touching the duodenal opening of the bile-duct with a glass rod dipped in dilute acid causes a gush of bile. The discharge of bile into the intestine is greatest about 3 to 5 hours and again at 13 to 15 hours after food, but it is subject to very considerable fluctuations.

The direct action of nerves upon pancreatic secretion, although probable, is not supported by definite experimental evidence. Stimulation of the medulla excites the secretion; excitation of the central end of any afferent nerve has a contrary effect. Destruction of the nerves that are distributed to the pancreas interferes with the above-mentioned effect, and is

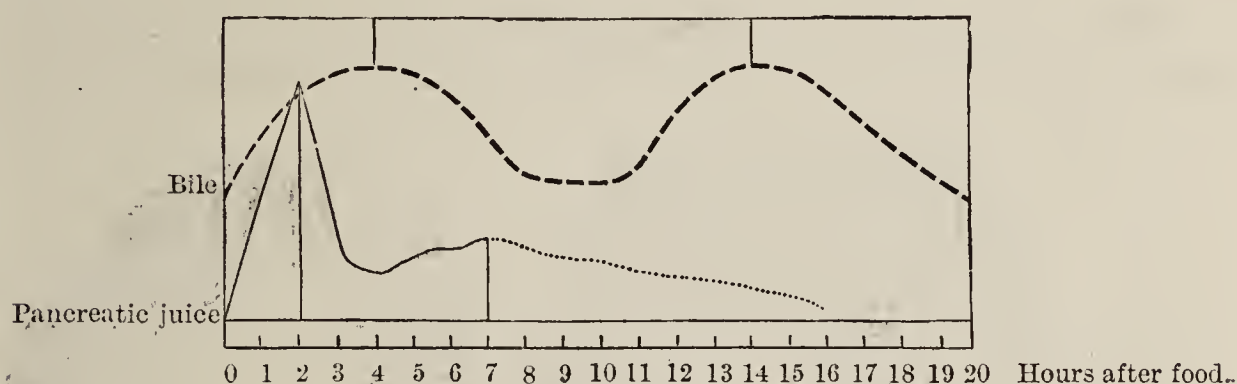


FIG. 86.

Curves of biliary and of pancreatic secretions subsequent to the ingestion of food.

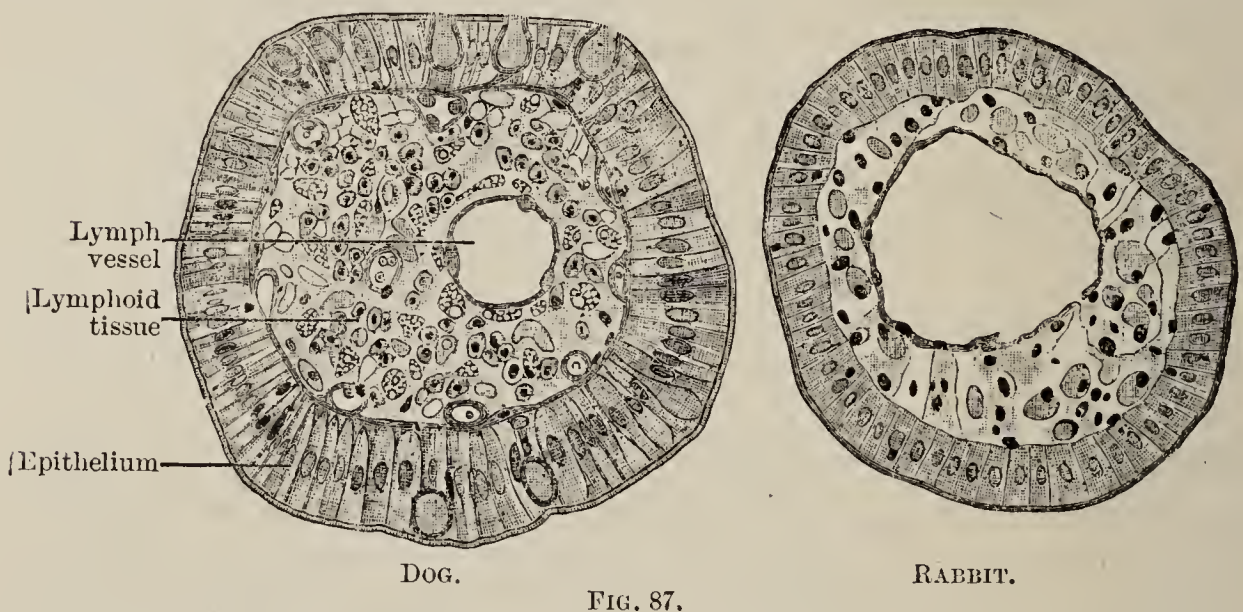
followed by the establishment of a thin watery secretion analogous with the 'paralytic' secretion that occurs from the sub-maxillary gland after section of the chorda tympani. But in the case of pancreatic, as in that of biliary secretion, the experimental isolation of secreto-motor nerves has entirely failed; we have no right to say that the increased and diminished secretion are independent of the increased and diminished vascularity with which they are always associated. We must be content to recognise in these cases the simple fact that increased secretion is associated with greater blood-supply, diminished secretion with smaller blood-supply, without attempting to lay down whether the associated changes bear a relation of cause and effect, or whether both are or are not the double effect of a common cause. Thus, with regard to any direct action of nerves upon pancreatic and biliary secretion, matters stand very much in the same position as in the case of gastric



secretion—viz. the existence of modifications of secretion by nervous influence is indicated as possible by pathological considerations as well as by certain rough results of experiments; for instance, the pancreatic and biliary secretions are accelerated in response to the stimulating effects of swallowed food, and, according to Heidenhain, stimulation of the spinal bulb distinctly augments the flow of pancreatic juice; but no undeniably direct secreto-motor action comparable with that of salivary nerves has ever been proved by the excitation of pancreatic, gastric, or hepatic nerves, and the possibility therefore remains that such effects as are observed are the indirect consequences of alterations of blood-supply.

### THE SMALL INTESTINE

The mass of chyme undergoing digestion is gradually propelled along the small intestine; this tube, if it were cut open



Transverse sections of villi of carnivorous and of non-carnivorous animals.  
(Heidenhain.)

and laid out flat, would occupy a large surface—about  $\frac{1}{2}$  square meter; and this already considerable surface is made still greater by the multitude of projections—*valvulae conniventes* and villi—with which it is beset; 10 square meters is a fair estimate of the total superficial area of the small intestine. The chyme is thinly spread—‘smeared’ over this large surface, a condition that is eminently favourable to absorption. *Absorption*, indeed, is the chief function of the small intestine, further digestion by intestinal fluid is its very subordinate function.

The wall of the small intestine is composed of the four coats

already alluded to, viz. mucosa, submucosa, muscularis, and serosa. From the physiological point of view, the most important feature to be considered is the minute anatomy of the villi, which constitute the organ of absorption; attention should also be paid (1) to the presence of Lieberkühn's follicles, which furnish the intestinal juice; (2) to the fact that lymphoid tissue enters largely into the composition of the mucous and submucous coats, especially in the ileum, where clusters of lymphoid nodules (Peyer's patches) are a prominent feature; and (3) to the fact that in the duodenum the submucous coat is occupied by a sheet of gland tissue (Brunner's glands), which is histologically identical with salivary gland. The physiological action of the nervous tissue that forms the plexus of Meissner in the submucous coat, and the plexus of Auerbach between the two muscular coats, is entirely conjectural.

The villi are the principal instruments of absorption. Their number has been estimated at 4 to 5 millions, and it has been calculated that each square centimeter of intestinal surface is increased twenty-fold by the villi which cover it. Their distribution is, however, far from being uniform, villi being longer, closer, and more numerous in the upper than in the lower portions of the small intestine. A villus is composed of epithelium and lymphoid tissue; it contains blood-vessels, lymphatics, and possibly nerves, also plain muscular tissue derived from the muscularis mucosæ; the amount of lymphoid tissue in a villus of a flesh-feeder (dog or cat) is much greater than in that of a vegetable-feeder (rabbit or guinea-pig).

By virtue of its muscular tissue a villus is contractile; in the relaxed state it is elongated, in the contracted state it is quite short, in consequence of the more or less stretched condition of the epithelial coat, according as the villus is relaxed or contracted; the individual cells are shorter and broader in a relaxed villus, longer and narrower in a contracted villus. According to Heidenhain, the adenoid tissue of the villi contains a large proportion of granular lymphoid cells peculiar in their behaviour to staining reagents and to osmic acid. They are blackened by the latter reagent, and it might therefore be supposed that the granular material in the cells is fatty; but that this is not so is shown by the fact that it is not dissolved out by ether. These

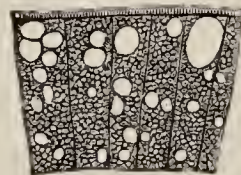


FIG. 88.

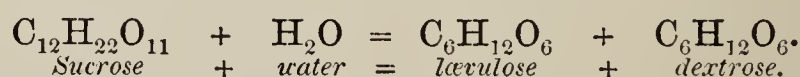
Intestinal epithelium taken during the absorption of fat, treated with ether and exhibiting vacuoles. (Heidenhain.)



cells are far more numerous during digestion than in the fasting state. They have been made responsible for the absorption of fat, but, as has just been stated, this is denied by Heidenhain; fat is absorbed by the epithelial cells themselves; that this is so, and that it does not pass between the cells nor is carried by the leucocytes, is shown by treating sections with ether, which dissolves the droplets of fat, leaving distinct vacuoles in the epithelial cells, while it has no action on the granular lymphoid cells that are stained by osmic acid (fig. 88).

The crypts of Lieberkühn are lined by epithelium, which at first sight appears identical with that covering the villi; they are not, however, on that account to be regarded as agents of absorption. They are in reality secreting glands. The epithelial cells of the crypts, as compared with those of the villi, stain more deeply and possess a much less distinct striated border; their nuclei frequently exhibit karyokinetic figures, which are hardly ever observed in the nuclei of cells lining the villi.

**Intestinal juice**, secreted by the glands of Lieberkühn (and by the glands of Brunner in the duodenum) is an alkaline fluid with a specific gravity of 1.010, having only a very slight digestive action upon the food-stuffs. Its most characteristic effect is the conversion of *cane-sugar* (sucrose) into *invert sugar* (lævulose + dextrose), by virtue of the ferment called *invertin*.



Intestinal juice has usually been obtained from an intestinal fistula—formed by cutting out from the tube a portion which is made to open externally, the cut ends of the main tube being brought together so as to restore its continuity. The isolated portion left in connection with its mesentery may be closed at one end, the other end being sutured to the abdominal wound; it then ultimately forms a blind sac with a single opening (Thiry's method), or both ends may be made to open at the surface of the abdomen (Vella's method).

An experiment by Moreau is quoted to show that nerves exercise some influence on the secretion of intestinal juice. A portion of intestine drawn out of the abdominal cavity is subdivided into three compartments by means of four equidistant ligatures; the nerves of the middle compartment are divided and the intestine replaced in the abdominal cavity. A few hours later the contents of the three compartments are examined; the middle

compartment is found to contain a considerable quantity of watery fluid, the two other compartments are almost empty. The effect, which may however be either a vascular or a direct secretory phenomenon, is comparable to that obtained on the salivary gland after section of the chorda tympani (paralytic saliva).

Before passing on to the consideration of intestinal absorption, opportunity may be taken to mention two important pathological relationships of intestinal glands. The glands of Brunner, of which we possess little or no physiological knowledge, are particularly prone to excessive action, going on to inflammation and to ulceration, in consequence of burns or scalds affecting any large extent of the cutaneous surface. The lymphoid patches of Peyer are particularly, it may even be said regularly, subject to inflammation and ulceration in the course of typhoid fever.

**Absorption** goes on in the whole intestinal tract—in the mouth, in the stomach, in the small and in the large intestine. It is at a maximum in the small intestine, in its upper portion more especially. Nearly everything that gets into the body does so by the surface of the upper part of the small intestine, after having undergone the digestive action of saliva, of gastric juice, of bile, and of pancreatic juice. We have already alluded to the fact that the conditions of absorption are (1) the diffusibility of the substance to be absorbed, (2) the physiological activity of the epithelium through which it is being absorbed. We have no right to say that either factor is the more important, but we have sufficiently realised the importance of the physical factor, and may therefore now insist upon the importance of the epithelial factor by adding to the illustrations already given of its efficacy (p. 162). Brücke long ago pointed out that egg-albumin can in part be absorbed as such, and his statement has found a practical application in the feeding of infants by raw white of egg, although we must now admit that in all probability the egg-albumin has become serum-albumin in the course of digestion. Voit and Bauer have determined that proteid solutions injected into the small intestine and into the rectum are actually absorbed, and the fact is no more than definite and exact proof that nutrient enemata are in reality nutrient. Bernard showed that if ferrocyanide and iodide of potassium be injected into a vein, the ferrocyanide passes into saliva and pancreatic juice, the iodide into the urine and bile; this is an illustration



of the selective activity of epithelium—not, indeed, in absorption, but in secretion—under identical physical conditions. Caze-neuve and Livon have shown that the living epithelium of the bladder forms a barrier to the absorption of urea. Reid, more recently, has given exact proof of epithelial activity in the absorption through frog's skin, by showing that the osmotic current, which is normally greatest from outer to inner surface of the living skin, is hastened by a stimulant (alcohol), retarded by a depressant (chloroform). We may sum up the function of the intestinal epithelium by saying that it furthers the absorption of favourable or feeding material, while it opposes the absorption of unfavourable or non-feeding material; its removal, so far from facilitating, would obstruct the absorption of food.

Observations on the rate of absorption are to the effect that about 1 cubic centimeter of fluid passes through 1 square centimeter of intestine per hour. Taking the villi into account, 1 sq. cm. of intestine has a surface of about 20 sq. cm., and the speed of the absorption-current through the epithelium will be about  $8\ \mu$  per minute, *i.e.* a given particle of fluid traverses the epithelial layer, which is 30 to 40  $\mu$  in thickness, in about 5 minutes.

We have sufficiently insisted upon the importance of the epithelium; with regard to lymphoid tissue, the part it plays in absorption is more obscure; it probably acts as a filter arresting the entrance of deleterious substances, or at least preventing their sudden entrance in mass, and giving time for their subsequent gradual elimination. In correspondence with this view we find a sheet of lymphoid tissue forming part of the absorbent surface, in the villi, in the submucous layer, and collected into nodules in the solitary and in Peyer's glands; we moreover find that sheet in its highest development in the large intestine, where the absorbent surface is most exposed to decomposition products; and finally, on the course of the larger lymphatics we find a second line of probable filtering organs, the mesenteric glands.

The channels of absorption are (1) the blood-vessels, (2) the lymph-vessels, with which, as we have seen, the villi are abundantly provided. The materials, the absorption of which we are to inquire into, are water, proteids, fats, carbohydrates, and salts.

The respective functions of the blood-vessels, and of the lymph-vessels, as channels of absorption, are, briefly, to the following effect: the greater part of the water, salts, sugar, and proteid absorbed passes into the blood-vessels, and by the portal

vein through the liver; the greater part of the fats absorbed passes into the chyle-vessels, and by the thoracic duct into the general venous system. In other words, the portal system is the chief channel of the absorption of water, salts, sugar, and proteid; the chyliferous system is the chief channel of the absorption of fats. We are nevertheless bound to admit that this distribution of function is not completely rigid: some small proportion of the water, salts, proteid, and sugar, does enter the chyle-vessels, some small proportion of fat does enter the portal system.

It is highly probable that the liver plays towards the portal blood a filter, or sentinel function, in arresting deleterious substances, like that played by lymphoid tissue and lymph-glands in relation to chyle. But this office is indicated only by pathological considerations, not by any definite experiments, and we did not therefore include it in the formal enumeration of liver functions.

We have now to follow as far as possible the internal digestion or the elaboration of the absorbed products of external digestion, and to answer the question—What happens to proteids, to carbohydrates, to fats, &c., when they have got into the body? We have studied the entrance of the proximate food-principles, and the preliminary transformations which they undergo; we can study their exit and the forms which they then assume, viz. carbon dioxide and urea; our present task is to study the chemical metamorphoses occurring between these two extremes. We shall find that with the exception of the glycogenic function of the liver, only the most fragmentary glimpses can be obtained of what takes place in the depths of the tissues.

### GLYCOGENESIS

The great importance of the glycogenic function of the liver is owing partly to the considerations just alluded to, partly to the fact that this function, while localised in and exclusively undertaken by the liver of fully-organised animals, is a function common to many tissues in the embryo, and essentially similar to the starch function in vegetable tissues. We shall therefore consider:—

(1) Its nature as manifested by the liver of an adult mammal. (2) Its distribution in the animal scale and in foetal tissues. (3) Its relation to food, health, and muscular action.

The glycogenic function of the liver was discovered by Claude



Bernard in 1848. He was making experiments to determine where sugar is consumed in the organism, and in the course of these experiments he analysed the blood going to and coming from the liver, and found that the blood of the hepatic vein contained more sugar than that of the portal vein; from which he concluded that the liver produces sugar. Thus, while looking for the seat of sugar-consumption, he found a seat of sugar-production. This was the first step. About ten years later Bernard supplemented his first discovery by finding that the liver contains a substance very like starch, which, like it, is convertible into sugar by ferment action, and which he therefore named 'glycogen.' This was the second step.

From these observations we learn that the liver can make sugar and can make glycogen, and recognise that the sugar-producing or glycogenic process consists of two stages:—(1) The formation of glycogen. (2) The conversion of glycogen into glucose.

*Objections.*—That the liver *normally* produces sugar was not at once accepted as unquestionable. Bernard himself had shown that an excised liver can produce sugar *post-mortem*; he showed that a liver washed sugar-free and left to itself is after a time found to contain a fresh supply of sugar. Pavy thereupon urged that the formation of sugar is not a normal physiological event, but only a *post-mortem* occurrence. He objected to Bernard's analyses of hepatic and of portal blood, that the hepatic blood had been allowed to stagnate in the liver, whose function was thereby disordered, and stated that normal hepatic blood contains no more sugar than normal portal blood. The case stood as follows:—According to Bernard, the liver is constantly forming sugar during life and continues to do so after excision; during life the sugar is carried off from the liver as fast as it is formed, so that a perfectly fresh liver is almost or quite sugar-free, while a stale liver contains an abundance of sugar. According to Pavy, it is not legitimate to conclude that the liver makes sugar normally in the body because it is found to do so out of the body. A fresh liver, pounded in a mortar with alcohol to remove any sugar present at the time, is then dried and powdered, and may be kept in this state for an indefinite time. This liver-powder, mixed with water and left for two or three hours in an incubator at 40°, gives an abundant sugar reaction. This experiment conclusively shows that a production of sugar can be effected in the liver-stuff, not merely

after the death of the animal, but also after the death of the liver itself.

The crucial point is the comparative analysis of portal and of hepatic blood. Do the two kinds of blood taken simultaneously from the two vessels without previous arrest of the hepatic circulation exhibit any difference in the amounts of sugar they respectively contain? Recent analyses show that they do, and that the average amount of sugar in portal venous blood is 1 per 1,000, in hepatic venous blood 2 per 1,000. The conclusion is therefore justified that the liver produces sugar in the body during its life, as well as out of the body after the circulation has ceased. If we take into account how great is the quantity of blood passing through the organ per hour or per day, we must recognise that the small difference between 1 and 2 per 1,000 is significant of a very large daily production of sugar. Seegen, for instance, has estimated that more than 400 litres of blood pass through the liver of a 40-kilo dog in one day, which, taking the sugar produced at 1 per 1,000, gives a daily production of over 400 grammes. This estimate is possibly an exaggerated one; still, there can be no doubt that the daily sugar-production by the liver is very considerable. With regard to the immediate antecedent of sugar, it is admitted by all authorities (with the exception of Seegen, who positively asserts that the liver can form sugar directly from either proteids, fats, or carbohydrates) that the liver-sugar is exclusively derived from glycogen.

*Preparation of glycogen.*—The preliminary condition is the choice of an animal with the liver in a highly glycogenic condition, *e.g.* a rabbit two or three hours after a large feed of carrots; or a dog two or three hours after a full meal of arrowroot. Moreover, having learned that glycogen is rapidly transformed into sugar *post-mortem*, we remove the liver *immediately* after death, and arrest the transformation by plunging the organ into boiling water, cutting or tearing it in pieces as quickly as possible. Under these conditions we shall find in solution a minimum quantity of sugar and a maximum quantity of glycogen. The boiling is continued for 15 to 20 minutes, the liver-tissue being further subdivided, and finally pounded with sand to ensure the extraction of all available glycogen. The mess is filtered and yields a milky opalescent solution of glycogen, mixed with some proteid and some fat. From this solution, when cool, proteid is removed by precipitation with Brücke's fluid (potassio-mercuric iodide) and dilute hydrochloric acid, added alternately in small quantities



until no further precipitate is produced. From the filtrate the glycogen is precipitated by the gradual addition of twice its volume of ordinary alcohol. The precipitate is collected upon a filter and washed, first with alcohol, then with ether to remove fat, finally with absolute alcohol.

The glycogen precipitate, dried over sulphuric acid, yields on pounding a bulky white amorphous powder, soluble in water, with which it gives an opalescent solution; this solution gives a red or brown colour with iodine, which disappears on heating and returns on cooling. As implied by its name, and by the synonym 'animal starch,' it is a carbohydrate body closely resembling starch and dextrin, convertible into the same product—sugar, and represented by the same empirical formula— $C_6H_{10}O_5$ .

*Source and fate of glycogen.*—We have now to consider the derivation of glycogen. Is it derived from proteids, or from fats, or from carbohydrates, or from all these three classes of foods? The answers to these questions are as follows :—

1. It is derived from food, because in the absence of food the liver contains no glycogen.

2. It is derived from carbohydrates especially, because animals fed upon exclusively starchy or saccharine food accumulate much glycogen in their livers. As regards the comparative effects of different kinds of sugar, Voit finds that dextrose and lævulose (which are most easily made to ferment by yeast) cause glycogen deposit in the liver whether they are absorbed from the intestine or directly injected into the circulation; that sucrose and maltose (which are less easily fermentable) produce this result only if they enter through the intestine; while lactose and galactose (which are hardly fermentable), whether they are swallowed or injected, do not give rise to any glycogen deposit.

3. It is also derived from proteids, because animals fed upon exclusively proteid food form glycogen in their livers. This, the cardinal fact of the whole debate concerning the source of glycogen, was for long the subject of dispute, and has only been determined by prolonged and repeated experiments. The observations were carried out upon dogs fed upon horse-flesh, and the objection that such meat contains glycogen is not of much weight, seeing that the amount in lean horse-flesh is less than 0·5 per cent.

4. It is not derived from fat, because animals fed upon an exclusive diet of fat lay by no glycogen in their livers.

Thus, as regards the formation of hepatic glycogen, we have

the following order of efficacy : carbohydrates most—proteids some—fats not at all.

The most important of these facts is that the nitrogenous proteids do give rise to the non-nitrogenous glycogen. It is strikingly exemplified in the pathology of diabetes, the essential symptom of which consists in the passage in the urine of a sugar with which the liver-sugar is identical. A subject of severe diabetes on ordinary diet (containing proteid, fat, and carbohydrate) passes an excessive amount of sugar, on a restricted diet (containing proteid and fat, but no carbohydrate) a smaller but yet a considerable amount of sugar, which must, therefore, be derived from proteid.

With regard to the influence of health and of exercise, the principal facts are that, *cæteris paribus*, an animal in good health, or a quiescent animal, or any hibernating animal, such as a frog at the outset of winter, has a liver with glycogen, while an animal in bad health, or an active animal, or an animal soon after emerging from hibernation, has a liver without glycogen. If, for instance, we look for glycogen and for sugar in the liver in the following cases, we should find :—

- |    |                                      |   |              |   |          |
|----|--------------------------------------|---|--------------|---|----------|
| 1. | In a liver from the post-mortem room | . | no glycogen  | . | no sugar |
| 2. | „ „ a butcher's stall                | . | no glycogen  | . | sugar    |
| 3. | „ „ a recently-killed tame rabbit    |   | glycogen and |   | sugar    |
| 4. | „ „ a hunted wild rabbit             | . | no glycogen  | . | no sugar |

These facts involve a consideration of the physiological relation between glycogen in the liver, sugar in the blood, chemical changes in the tissues. Sugar is a constant constituent of the blood, which in a starving or a well-nourished animal alike contains about 1 per 1,000 of sugar. The amount of sugar present in the venous blood of muscle is constantly slightly below that present in arterial blood, and the amount present is at its minimum in the venous blood of active muscle.<sup>1</sup> Sugar is therefore consumed by living muscle, especially during activity. The amount of sugar present in hepatic venous blood is constantly slightly above that present in portal venous blood. Sugar is therefore produced by the liver. We have now no difficulty in following the sugar-cycle in an active well-fed animal. Absorbed sugar enters the blood, is to a great extent stored as glycogen in

<sup>1</sup> Chauveau estimated, *e.g.*, that 175 grms. of blood traversed 1 kilo of resting muscle in 1 minute, losing 0.036 gm. sugar, whereas through active muscle the quantities per minute were 850 grms. and 0.140 gm.



the liver, is consumed by living muscle, is discharged as  $\text{CO}_2$  and as  $\text{H}_2\text{O}$ . The part played by glycogen is that of a temporary carbohydrate reserve.

In a starving animal, receiving no fresh supply of sugar, sugar is nevertheless a necessity; the glycogen store soon disappears; yet the animal lives on for a time, having sugar in its blood, and using up sugar in its muscles. It is probable that this sugar is made *by* the tissues *from* the tissues (by the liver and possibly by muscle, from proteid and possibly from fat). From this point of view the fundamental chemical event is the formation and consumption of sugar; the formation of glycogen is a first incident in this event, occurring whenever the carbohydrate supply is plentiful, and failing to occur when the food-supply is suspended.

It is instructive to observe that this reserve carbohydrate function is represented throughout the animal and vegetable kingdoms; the liver makes and puts by glycogen at one time, transforms and discharges glycogen at another time; a potato makes and puts by starch at one time, transforms and discharges starch at another time. The alternate phases are of very different duration in the different cases; the carbohydrate reserve remains for a few hours or days in the liver of a warm-blooded animal, for many days or weeks in the liver of a cold-blooded animal, for several months in the tuber of a potato, or in the liver of a hibernating animal.

*Alterations of the liver structure* in consequence of digestion are obvious enough both to the naked eye and microscopically, but the interpretation of their significance in all their visible details is rendered uncertain, mainly because we have to deal with two possible altering agents—with an outgoing current of secretion as in other glands, and with an incoming current of food-products. It is to the last-named factor that the more obvious changes are due. The liver of a rabbit dieted in preparation for glycogen extraction is large, pale, and twice as heavy as the liver of a similar rabbit not so prepared; and whereas the liver of the fed rabbit may yield over 10 grammes of glycogen, that of the other rabbit would yield less than 1 gramme. A microscopic section of a dog's liver during the fasting state shows uniformly granular shrunken cells with obscured nuclei; taken twelve or fourteen hours after a copious meal, the cells are distended with coarse clumps, which give a red-brown colour on irrigation with iodine, and which, when they dissolve, leave the

cells with distinct boundaries and distinct nuclei in a ragged intracellular network of protoplasm. The appearance is evidently due to a deposition of and distension by glycogen during the post-prandial period.

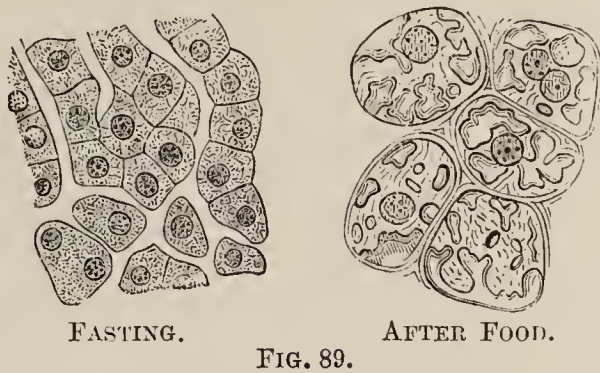
Changes in the main similar to the above and attributed to the glycogenic process have also been made out upon frogs, in which the liver is of distinctly tubular type. According to

Langley, the liver of a winter frog exhibits a distinction into zones similar to those of the

salivary glands; in this case the outer hyaline zone is loaded with glycogen, the inner granular zone with material presumably proteid, but which, seeing that the liver yields no definite ferment, cannot be termed 'zymogen.'

Another material that is apt to be deposited in the liver-cells, both of mammalia and of frogs, is fat, which forms droplets more or less abundantly in accordance with the state of nutrition (*e.g.* the fatty livers of crammed geese or of overfed and ill-nourished persons), but in no causal relation with the particular phase of digestion.

Finally, we have to recognise that the hæmolytic function of the liver may be attended with visible changes. According to Delépine, a granular deposit giving the ferric reaction exists in the liver-cells—slightly during fasting, not at all immediately after a meal, most abundantly eight to twelve hours later; and whereas the glycogen deposit is most abundant around hepatic veins, the iron deposit is thickest round the portal veins.



Liver-cells of dog after a thirty-six hours' fast, and fourteen hours after a full meal—in the latter case swollen with glycogen. (Heidenhain.)



## CHAPTER VI

### *RENAL EXCRETION*

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**The relation of excretion to digestion.**—Respiration, digestion, circulation, and excretion are the agencies through which the blood is maintained of such quality and composition that it serves throughout life as the vehicle of nourishment and of purification to the whole body. Respiration is essentially a process of exchange, formed by a double current of gases—of incoming oxygen, and of outgoing carbon dioxide. Digestion and excretion similarly constitute a process of exchange, in which, however, the two currents run in separate channels; carbon and nitrogen entering the body together by intestinal absorption, but nitrogen leaving the body by renal excretion as urea, and carbon by respiratory excretion as  $\text{CO}_2$ . And whereas in the absorption of oxygen the gas exists ready in the air, and enters the body unaltered, carbon and nitrogen are conveyed into the body in

certain combinations or proximate principles (carbohydrates, fats, and proteids) which form part of our ordinary articles of diet, and require to be prepared for absorption by digestion in the alimentary canal before they can actually enter the body. Thus respiration is simple absorption and excretion of gases, digestion is the preparation for absorption, and actual absorption of proteids, of carbohydrates, and of fats. Indigestible substances simply pass through the alimentary canal without actually entering the body, and are got rid of by defæcation, in company with an insignificant amount of matter actually excreted by the intestinal canal. The principal excretions, properly so called, are  $\text{CO}_2$  by respiration, and urea by renal action. We have already considered the former, we are about to consider the latter.

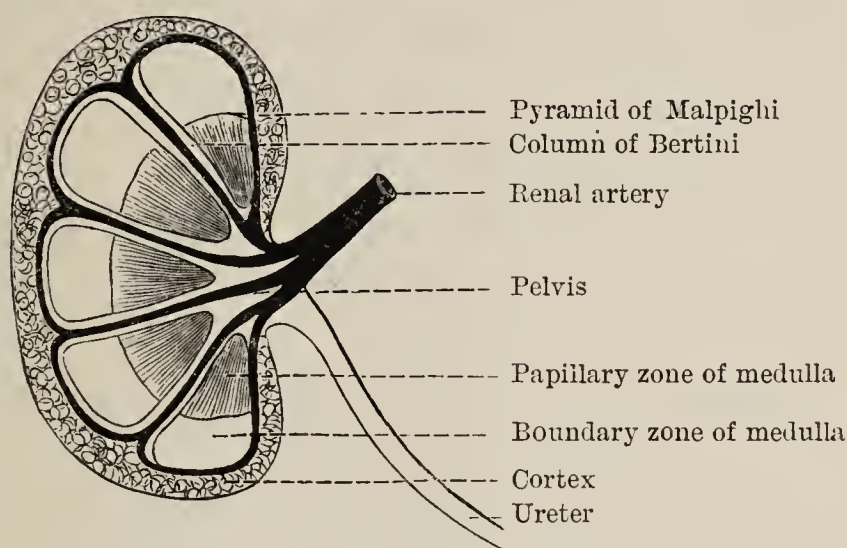


FIG. 90.—DIAGRAMMATIC SECTION THROUGH THE KIDNEY.

The branches of the renal artery pass along the columns of Bertini and form a series of arterial arches between cortex and medulla; from these arches spring the interlobular arteries and the vasa recta. (See fig. 92.)

**The kidney.**—A human kidney sliced longitudinally exhibits an outer zone—the cortex—surrounding an inner portion—the medulla. The medulla is composed of twelve to fifteen pyramidal masses; its outer portion in contiguity with the cortex is called the boundary zone; its inner portion is termed the papillary zone, each pyramid terminating as a papilla which protrudes into the pelvis of the kidney; the pelvis is the common central cavity and the commencement of the ureter. The structural characters of the several portions of the kidney are determined by peculiarities in the anatomical disposition of (1) the blood-vessels, (2) the urinary tubules. Microscopically, the cortex is characterised by convoluted tubules in such mass as to be termed ‘the labyrinth’; it also contains bundles of straight tubules forming the medullary rays. The boundary zone of the medulla is



characterised by looped tubules, the papillary zone by converging tubules. In an injected kidney the cortex is characterised by the presence of capillary 'glomeruli,' the boundary zone by the presence of a straight system of capillaries; the cortex contains also a rich network of capillaries around the convoluted tubes; the papillary zone of the medulla is, on the contrary, very scantily supplied by capillaries, which are derived from the vasa recta. As in all secreting or excreting glands, the essential elements are a thin sheet of *blood* separated by a *membrane* from a layer of *epithelial cells*. In the case of the kidney the blood is contained in the capillaries forming the glomeruli and surrounding the tubules, the epithelium covers the glomeruli and lines the tubules.

The renal artery enters at the hilum and divides at once into several main branches, which run between the pyramids

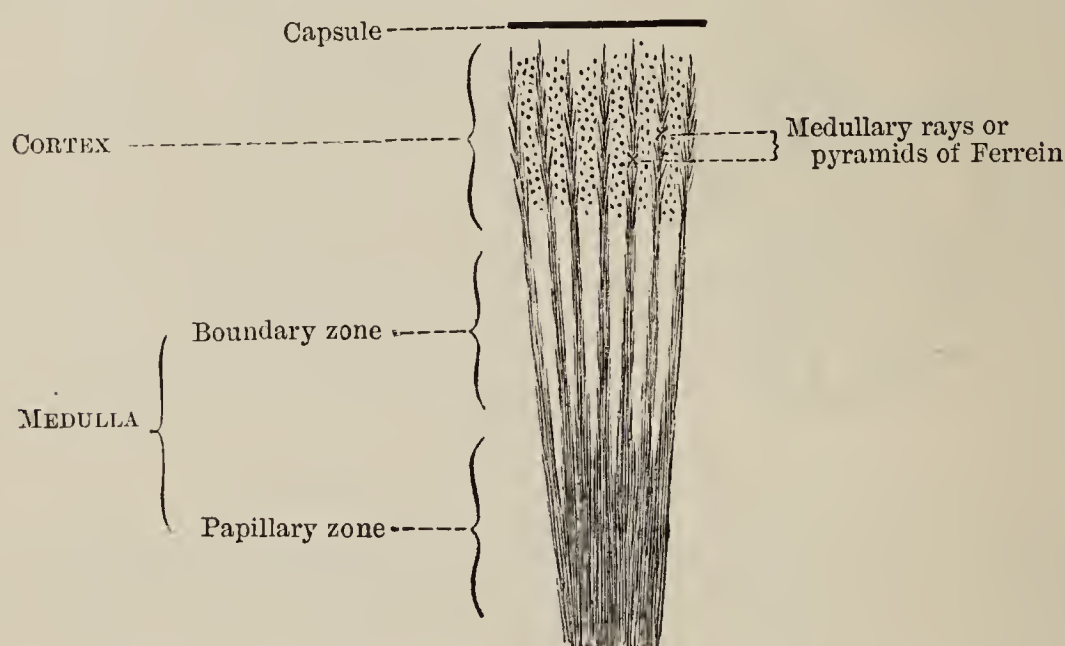


FIG. 91.—DIAGRAMMATIC SECTION THROUGH A PORTION OF THE KIDNEY.

towards the cortex; at the junction between medulla and cortex the branches form incomplete arcades, from the two sides of which spring the smaller vessels supplying the labyrinth (interlobular vessels) and the boundary zone (vasa recta). From each side of an interlobular artery spring several short branches (afferent vessels), each of which breaks into a tangled bunch of capillaries constituting a glomerulus; the minute veins (efferent vessels) formed from these capillaries do not at once unite with other veins, but first undergo redistribution as a second capillary network surrounding the convoluted tubes; some of these efferent vessels, springing from glomeruli close to the boundary zone, form a second capillary network in that zone (false vasa recta). The peculiarity of the renal circulation consists in this

double capillary network—a first capillary distribution forming the glomeruli, a second capillary distribution, after the manner of a portal vein, forming the vascular network over the tubules. The disposition is similar in principle, though not in detail, with that found in the kidneys of amphibia, where glomeruli and tubules are supplied by distinct vessels—glomeruli by the renal artery, tubules by the renal portal vein (which is itself a branch of the femoral vein).

A urinary tubule commences by a capsule (Bowman) surrounding and reflected over the glomerulus. From this com-

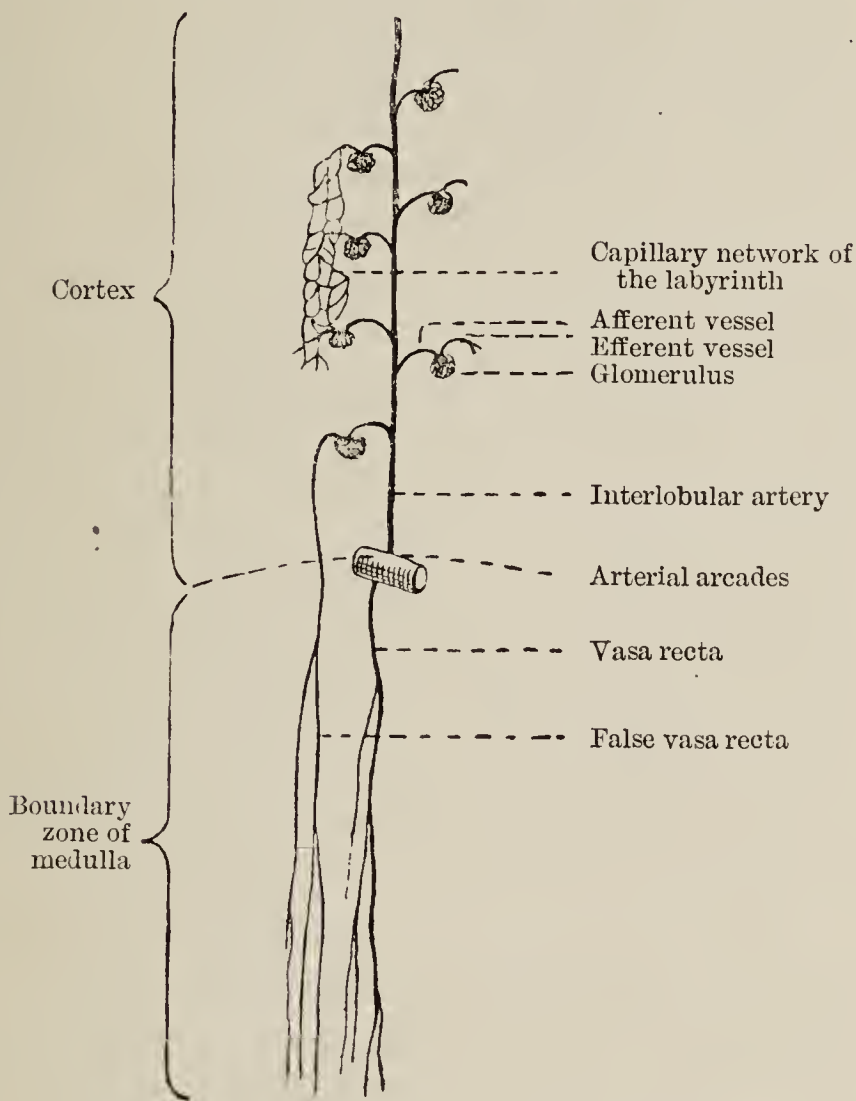


FIG. 92.—DISTRIBUTION OF RENAL ARTERIES.

mencement to its end at a papilla it runs a very complex course, and is lined by epithelium of very varying appearance. Omitting details of purely morphological interest, the course and character of a tubule are as follows:—The glomerular capsule is composed of two layers, the inner of which covers and is fused with the glomerular loops, while the outer layer is continued as the tube itself. This tube while in the cortex is convoluted, in the next portion of its course it forms a loop (Henle) occupying the whole breadth of the boundary zone of the medulla; in this loop we



have to distinguish physiologically, as well as morphologically, the broad ascending from the narrow descending limb; re-entering the cortex, the tube again becomes convoluted until

it joins one of the straight tubes in the medullary ray; from this point onwards the tubes run straight down the medullary ray and pyramid, joining tube with tube, until they terminate by orifices on the papilla. The epithelium lining this tract is distinguishable into three varieties: (a) Squamous epithelium lining the capsule. (b) A more or less granular and striated glandular epithelium lining the convoluted portions in the cortex, and the ascending limb of the loop in the boundary zone. (c) A clear cubical lining epithelium in the descending limb of the loop, and in the straight tubes running in the medullary ray and in the pyramid. Of these three epithelia, the second is of most physiological importance; its individual cells are active masses of protoplasm concerned in extracting from the blood the chief organic constituents of urine, and, upon occasion, certain foreign materials experimentally injected.

**The urine.**—The urine is continuously secreted by the kidneys and passed along the ureters to the bladder, where it accumu-

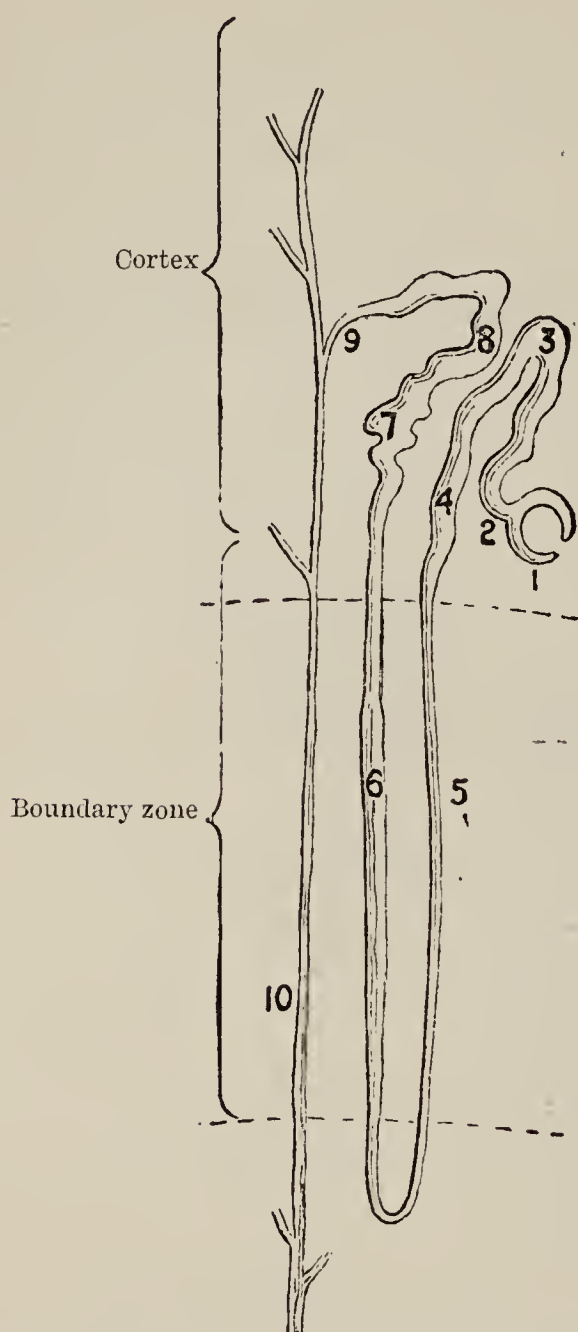


FIG. 93.—COURSE OF A URINARY TUBULE.

1. Capsule of Bowman.
2. Neck.
3. First convoluted tube.
4. Spinal tube.
5. Descending limb of loop of Henle.
6. Ascending limb of loop of Henle.
7. Irregular tube.
8. Second convoluted tube.
9. Junctional tube.
10. Collecting tube.

lates, and from which it is discharged at intervals by the act of micturition. The average amount of urine secreted per diem is 1,500 c.c., or about 50 oz., and the capacity of the bladder is about 500 c.c.

*Composition of the urine.*—Fresh normal urine is a clear yellow

fluid, of acid reaction, with a specific gravity of 1,015 to 1,025, being a solution in water of *urea* and *salts*, viz. about 2 per cent. of urea and  $1\frac{1}{2}$  per cent. of salts. Stated more in detail, the composition of urine as regards its most important constituents, and the average daily discharge of such constituents, are as follows:—

	per 1,000	per diem	
Water . . . . .	960 . . . . .	1,440	c.cm.
Urea . . . . .	20 . . . . .	30	grammes
Uric acid . . . . .	2 . . . . .	0.75	„
Hippuric acid . . . . .		0.75	„
Creatinin . . . . .		1.5	„
Phosphates . . . . .	15 . . . . .	H <sub>3</sub> PO <sub>4</sub> . . . . .	3 „
Chlorides . . . . .		HCl . . . . .	7.5 „
Sulphates . . . . .		H <sub>2</sub> SO <sub>4</sub> . . . . .	3 „
&c., <i>e.g.</i> mucus and extractives	3		

The *acidity* of urine is not due to the presence of any free acid; methyl orange, which is reddened by free acid, is not affected by urine; sodium hyposulphite, which with free acid gives a precipitate of sulphur, gives none with urine. The acidity is attributable to the presence of acid sodium phosphate, NaH<sub>2</sub>PO<sub>4</sub>, and of free carbonic acid gas. Normally the acidity of fresh urine increases a little at first; it subsequently diminishes and is replaced by alkalinity, this last change being due to a fermentation, in the course of which urea is converted into ammonium carbonate. The normal acidity of urine is equivalent to that of a 1 per 1,000 solution of H<sub>2</sub>SO<sub>4</sub>. The acidity is somewhat more pronounced in the urine passed in the morning, less so in that passed after a meal, and it varies greatly with the nature of the diet; the urine of herbivorous animals and of vegetarians is alkaline and turbid, that of carnivorous animals and of ordinary men is acid and clear, that of herbivorous animals during starvation and during hibernation—when they are practically carnivorous of their own flesh—is also acid and clear. Abnormally the urine may become excessively acid or alkaline. In the first case, a ‘cayenne-pepper’ deposit of uric acid (visible to the naked eye), and ‘envelope crystals’ of oxalate of lime, are discoverable by the microscope in the urine that has been allowed to stand. The second state may be due to disease of the kidney or bladder, causing the urine to decompose before it is discharged, or it may be caused by nothing more serious than an excessive consumption of fruit or vegetables; ‘knife-rester’ crystals of ‘triple phosphate’ (MgNH<sub>4</sub>PO<sub>4</sub>) and ‘hedge-



hog crystals' of ammonium urate are apt to form in such urine.

A normal clear urine may become turbid on cooling by deposition of urates; on the other hand, a normal urine may become turbid on heating by deposition of phosphates; in the first case the turbidity is removed by heating, in the second it is removed by acetic or nitric acid.

The *specific gravity*—1,015 to 1,025—varies with the amount of urine passed; this again varies with the amount of water discharged from the lungs and skin. Thus, in winter the urine is dilute, pale, and copious, as compared with the summer urine, which is concentrated, dark, and scanty. Normally the total amount of solids per 1,000 contained in a sample of urine is approximately indicated by multiplying the last two figures of

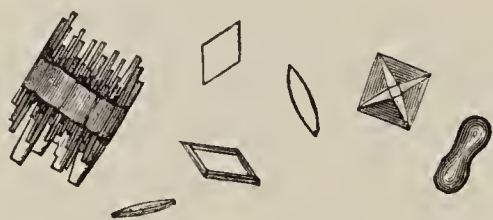


FIG. 94.—CRYSTALS IN ACID URINE;  
URIC ACID AND OXALATE OF LIME.

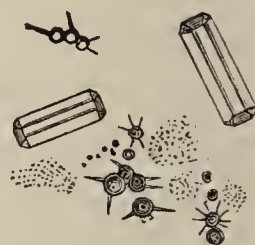


FIG. 95.—CRYSTALS IN ALKALINE URINE;  
TRIPLE PHOSPHATE AND AMMONIUM URATE.

the specific gravity by 2·2; *e.g.* with S.G. 1,015 the total solids per 1,000 are 33; with S.G. at 1,020 they are 44 per 1,000.

The *colour* of the urine is due to the presence of urobilin, which, as indicated by its name, is a derivative of bile-pigment; urobilin is, in fact, identical with hydrobilirubin.

**Mechanism of secretion; theories.**—The secretion of urine is an active process, not a passive filtration. On the passive filtration theory (Ludwig, 1844 to 1870), it was supposed that the entire urine is filtered under pressure from the blood in a very dilute state at the glomeruli, and that it gradually becomes more and more concentrated by re-absorption of water as it passes along the tubules. On the active secretion theory (Bowman 1842, Heidenhain 1875), it is supposed that the epithelial cells lining the glomeruli and the convoluted tubules separate the water, urea, and other urinary constituents from the blood, and that different shares of work belong to the two classes of cells—water being separated by the cells of the glomeruli; urea, uric acid, &c. by the cells of the convoluted tubes and of the ascending limb of Henle's loop. The anatomical disposition of the renal vessels is such as to suggest filtration under high pressure at the glomeruli,

secretion under low pressure by the convoluted tubules; and in favour of the filtration theory we have also the fact that, as a rule, the amount of urine is greater and smaller as blood-pressure rises and falls. But, as we shall see, there are exceptions to the rule, and the facts can be otherwise explained. The active secretion theory is upheld by the manner in which the secretion of indigo carmine has been traced by Heidenhain, and is supported by Nüssbaum's experiments on the kidneys of amphibia; it is not contradicted by any of the facts advanced in support of the pressure theory.

Moreover, a simple comparison of the percentages of substances contained in the blood and in the urine respectively is highly suggestive of separate activity rather than of passive filtration; the normal percentage of urea is 2 per cent. in urine and only  $\cdot 02$  per cent. in normal blood, and not above  $\cdot 2$  per cent.



FIG. 96.—RABBIT'S KIDNEY.

One hour after division of spinal cord and injection of 5 c.c. sulphindigotate of soda. (Heidenhain.)



FIG. 97.—MICROSCOPIC APPEARANCE OF CONVOLUTED TUBULES: (Heidenhain.)

in extreme uræmia; in diabetes the percentage of sugar in the urine has been found to reach to ten times that in the blood.

Heidenhain's experiments with indigo carmine are as follows:—The spinal cord or bulb of a rabbit is divided in order to lower the blood-pressure, indigo carmine (5 to 20 c.c. of a saturated solution of sodium sulphindigotate) is injected, the animal is killed ten minutes to an hour later, its kidneys are removed and injected with alcohol. On slicing open the kidney, which has become blue, the coloration is seen in a well-marked band which occupies the whole breadth of the cortex, and sometimes in a fainter band corresponding with the boundary zone; on microscopic examination the coloration is found to be due to blue granules in or between the epithelial cells—(1) of the convoluted tubes, (2) of the ascending limb of Henle's loop, and (3) in



the lumen of the tubes themselves; the capsules of the glomeruli are found to be quite free from blue granules. This typical experiment is accepted as an illustration by means of a visible substance of what probably is the mode of separation of the invisible substance, urea. The division of the spinal cord has prevented any copious stream of water from flowing through the glomeruli, and the blue granules have remained in or near the parts engaged in their separation.

Various modifications of the above typical experiment are taken to be confirmatory of this interpretation. The indigo injection has been made without previous division of the spinal cord, and the blue granules have been found massed in the pyramids or in the pelvis, as if they had been washed onwards by the water-stream. This last experiment has been repeated with previous corrosion of a superficial portion of cortex by silver nitrate, so as to interfere with the action of the subjacent glomeruli, and the wedge of kidney below the cauterised surface has been found with blue zones, as if unwashed, contrasting with adjacent parts in which the water-current had proceeded unchecked.

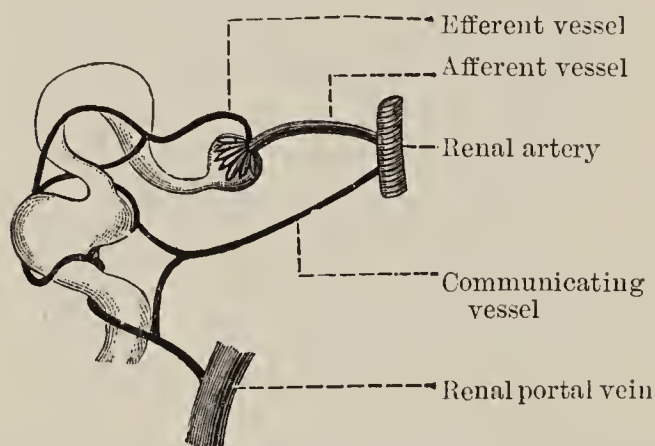


FIG. 98.—DIAGRAM TO ILLUSTRATE THE DOUBLE BLOOD-SUPPLY OF THE NEWT'S KIDNEY. (After Nüssbaum.)

The renal artery forms the glomeruli; the renal portal vein supplies the capillaries of the tubules.

The two systems of capillaries are not completely distinct; the connection is such that while the glomeruli are supplied chiefly by the renal artery, the tubules are supplied chiefly by the renal portal vein.

These results are consonant with the view that water is secreted by the glomeruli, urea (as represented by indigo carmine) by the tubuli contorti.

In prosecution of these inquiries, Nüssbaum utilised the amphibian kidney, which presents certain advantages for experimental purposes, by virtue of its double blood-supply from the renal artery and from the renal portal vein—the artery mainly supplying the glomeruli, the vein mainly supplying the tubuli. Nüssbaum's most pertinent observation was to the effect

that the sulphindigotate is discoverable in the tubuli after ligature of the renal artery; he did not, however, perform the counter-experiment of ligature of the vein. He found that albumin, sugar, carmine, do not pass into the urine after ligature of the renal artery, and concluded that these substances were secreted by the

glomeruli. He also found, however, that by injecting urea he could get secretion of water containing urea, even after ligature of the renal artery. This contribution of Nüssbaum's gives very little direct support to the theory of Heidenhain, and the method is open to the objection that ligature of the renal artery does *not* arrest circulation through the glomeruli.

*Effects of blood-pressure.*—The secretion of urine rises and falls with rising and falling blood-pressure; clinically, in cases of heart-disease, a falling-off in the amount of urine gives warning that the circulation is failing. But that the secretion is not a simple consequence of the pressure of the blood in the renal vessels is shown by the effect of compressing the renal vein, which, while not lowering the pressure, arrests the secretion. A more correct and comprehensive general statement is that the secretion of urine increases and diminishes with increased and diminished *blood-flow*. A justification of this statement is to be found in the subjoined table, which summarises the effects of various kinds of experiments upon the blood-pressure in the renal vessels, upon the blood-flow through the kidney, and upon the amount of urine secreted. More or less urine is given out according as more or less blood comes under the influence of the secreting cell, and if rising and falling blood-pressure produce as a rule more or less secretion, it is because rising and falling blood-pressure generally entail increased and diminished blood-flow.

—	Renal blood-pressure	Renal blood-flow	Secretion of urine	Volume of kidney
Destruction of spinal bulb . .	<i>dimin.</i>	<i>dimin.</i>	<i>dimin.</i>	—
Excitation „ . .	<i>incr.</i>	<i>dimin.</i>	<i>dimin.</i>	—
Section of renal nerves . .	—	<i>incr.</i>	<i>incr.</i>	<i>incr.</i>
Excitation „ „ . .	—	<i>dimin.</i>	<i>dimin.</i>	<i>dimin.</i>
Excitation of spinal bulb after section of renal nerves . .	<i>incr.</i>	<i>incr.</i>	<i>incr.</i>	<i>incr.</i>
Compression of renal artery . .	<i>dimin.</i>	<i>dimin.</i>	<i>dimin.</i>	<i>dimin.</i>
„ „ renal vein . .	<i>incr.</i>	<i>dimin.</i>	<i>dimin.</i>	<i>incr.</i>
Failure of heart . . . .	<i>dimin.</i>	<i>dimin.</i>	<i>dimin.</i>	—
Digitalis . . . . .	<i>incr.</i>	<i>incr.</i>	<i>incr.</i>	—

These several effects are the simple physical consequences of varying vaso-constriction throughout the system and in the kidney. It is hardly necessary to explain them at length; the facts are stated in the table in their simplest form. Destruction



of the bulb relaxes the vessels all over the body, inclusive of the kidney, and the circulation is everywhere more sluggish. Excitation of the bulb constricts the vessels all over the body, inclusive of the kidney, and although pressure is raised, less blood gets through the constricted vessels of the kidney; if, however, these vessels be cut off from vasomotor action by section of the renal nerves, then excitation of the bulb has the effect of sending a larger amount of blood through the kidney.

The method by which the circulation of the kidney can be most directly studied is to observe the alterations of volume which the organ undergoes. An instrument devised by Roy for this purpose, termed the oncograph, has already been described (p. 78). It consists of two parts united by a tube, one the exploring chamber or 'oncometer,' which receives the kidney, the other the recording chamber or 'oncograph' proper, which carries a recording lever fixed to a piston. The apparatus being filled with oil, any swelling or shrinking of the kidney causes the piston to rise or fall. Variations of blood-pressure cause corresponding variations of volume; the first column of the table, therefore, holds good for the alterations of volume produced by various experiments; *i.e.* the kidney shrinks with vaso-constriction (rise of pressure), swells with vaso-dilatation (fall of pressure); if, however, its nerves are divided, then it swells with rise of pressure and shrinks with fall of pressure. In reasoning out the connection between pressure and volume, it is to be remembered that the renal arteries share in the general contraction which causes rise of pressure, therefore the kidney shrinks with rise of pressure by vaso-constriction, but that they are also dilatable; therefore, unless the kidney shrinks by vaso-constriction, it passively swells from any increase of pressure, *e.g.* by respiratory undulations of pressure, or by excitation of the bulb after section of renal nerves. The only case in which there is anything to add is that of section and excitation of the renal nerves, of which we could not definitely state the effects on renal pressure, but which very obviously produce—the first, a dilatation, the second, a contraction of kidney volume.

We may take this occasion to define the physiological character of the renal nerves. Experiments by section and excitation demonstrate clearly the existence of vaso-constrictor action, less clearly that of vaso-dilatator action, not at all that of secretomotor action. That constricting fibres exist in the renal nerves

is shown by expansion of the kidney after the nerves are cut, and by the contraction of the kidney which is the usual result of their excitation; that dilating fibres also exist in the renal nerves is shown by the expansion of the kidney which is the occasional effect, or after-effect, of such excitation (Bradford), but, as in other nerves, the dilating are ordinarily masked by constricting effects. Renal nerve-fibres (in the dog) leave the spinal cord by the anterior roots from sixth thoracic to second lumbar, in greatest abundance by the last three or four thoracics. In the cord itself, vasomotor influence to the kidney passes down *both* sides, but most down the same side.

**Origin and separation of urea.**—The physiological history of urea as known to us at the present day may be summed up as follows:—(1) *Urea is the chief product of the disintegration of proteid food*; (2) *it is formed in the tissues*; (3) *it is discharged into the blood*; (4) *it is separated from the blood by the action of the renal epithelium*; (5) *it is expelled in the urine*. The first of these propositions is proved by the series of experiments in course of which the old view, that urea is the measure of muscular waste, was disproved, and replaced by the modern view, that urea discharged is in proportion with proteid food consumed. These observations, as the outcome of which the doctrine of nitrogenous equilibrium was formulated, will be considered in Chapter VII, and we shall at this stage confine ourselves to the recognition of the fact that urea is mainly derived from proteid which has not actually become protoplasm, but which has become disintegrated under the influence of protoplasm. The simplest and most cogent proof of the second, third, and fourth propositions is furnished by the fact that urea continues to be formed and accumulates in the blood (uræmia) when renal secretion has been abolished by removal of the kidneys, or by disease, and we shall see later that the liver is to be recognised as the most prominent among urea-making tissues (p. 242); specific proof of the separative activity of renal epithelium is afforded by Heidenhain's observations of the excretion of indigo carmine and of uric acid. The last proposition needs no formally stated proof; we know that urea is contained in the urine.

**Urea** is the chief normal constituent of urine, and the vehicle in which nitrogen is discharged from the body. It is a crystalline body soluble in water and in alcohol, capable of forming compounds with nitric acid (nitrate of urea), with oxalic acid (oxalate of urea), and with mercuric nitrate, and of being decomposed by fermentation and by



solutions of the hypobromites or hypochlorites. Urea has the empirical formula  $\text{CON}_2\text{H}_4$ , its molecular weight is  $12 + 16 + 28 + 4$ , or 60, of which nitrogen constitutes 28 parts, or nearly one-half. Thus, 60 grammes of urea will yield 28 grammes or 22.38 litres of nitrogen; 1 gramme of urea in urine should yield 373 c.c. of nitrogen at normal temperature and pressure, but is actually found to yield only 354 c.c., so that 1 c.c. N represents  $\frac{1}{3\frac{1}{5}\frac{1}{4}}$  gramme urea. The rational formula of urea is  $\text{CO} \cdot 2(\text{NH}_2)$ , and exhibits it as the diamide of carboxyl. It is isomeric with ammonium cyanate,  $\text{NH}_4\text{-O-CN}$ , and as a first stage in its alkaline fermentation yields ammonium carbonate,  $\text{CO}(\text{NH}_2)_2 + 2\text{H}_2\text{O} = (\text{NH}_4)_2\text{CO}_3$ . By further fermentation it decomposes into  $\text{CO}_2$  and N, and the same result is obtained in consequence



FIG. 99.—UREA. (From Funke's Atlas.)

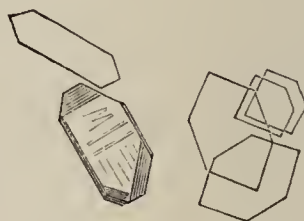
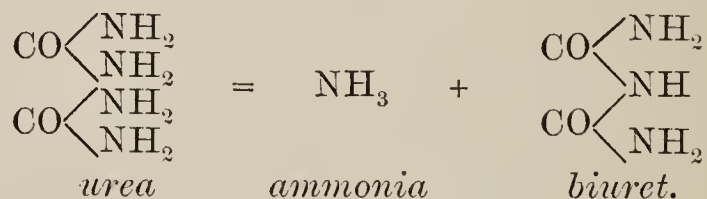


FIG. 100.—NITRATE AND OXALATE OF UREA.

of decomposition by chemical reagents; facts which furnish means for estimating a *quantity* of urea by a *volume* of nitrogen evolved.

Crystals of urea, heated in a dry test-tube to  $150\text{--}160^\circ$ , melt and give off ammonia; the residue is biuret, which if dissolved in water and tested by a trace of  $\text{CuSO}_4$  and excess  $\text{KHO}$ , exhibits a violet colour; if heated above  $170^\circ$ , either urea or biuret will yield ammonia and cyanuric acid.



*Synthesis of urea.*—Chemically as well as physiologically urea is of great importance as being the first animal product that has been reproduced by synthesis from inorganic bodies, or from the elements themselves, without the intervention of animals. Nitrogen passed through a tube containing carbon and potassium carbonate heated to redness gives potassium cyanide;  $\text{K}_2\text{CO}_3 + \text{N}_2 + 2\text{C}_2 = 2\text{KCN} + 3\text{CO}$ ; by oxidation the cyanide gives cyanate; potassium cyanate with ammonium sulphate gives potassium sulphate and ammonium cyanate;  $2\text{KCNO} + (\text{NH}_4)_2\text{SO}_4 = \text{K}_2\text{SO}_4 + 2\text{NH}_4\text{CNO}$ ; evaporation of ammonium cyanate yields urea.

*Preparation of urea from urine.*—Evaporate one or two litres of urine on a water-bath to a small bulk. Add nitric acid to form nitrate of urea. Pour off the fluid, and mix the crystalline mass with barium carbonate. Extract the mixture with alcohol. Filter, evaporate, and dry the crystalline residue of urea. Purify by recrystallisation.

The *presence of urea* is recognised in a fluid by the crystals of nitrate of urea that are formed when a drop of the fluid to which nitric acid has been added is allowed to evaporate on a slide and examined by the microscope. Nitrate of urea appears in the form of six-sided rhombic tables. The presence of urea may be suspected in a fluid that gives out bubbles on the addition of hypobromite of soda.

The *quantity of urea* in a fluid known to contain urea, *i.e.* in urine, is most conveniently and rapidly ascertained by the hypobromite method (Knop-Hüfner). It should first be remarked that an estimation of urea in a single sample of urine is of little value; the estimation, to be of any use, must be taken with a sample of the entire urine col-

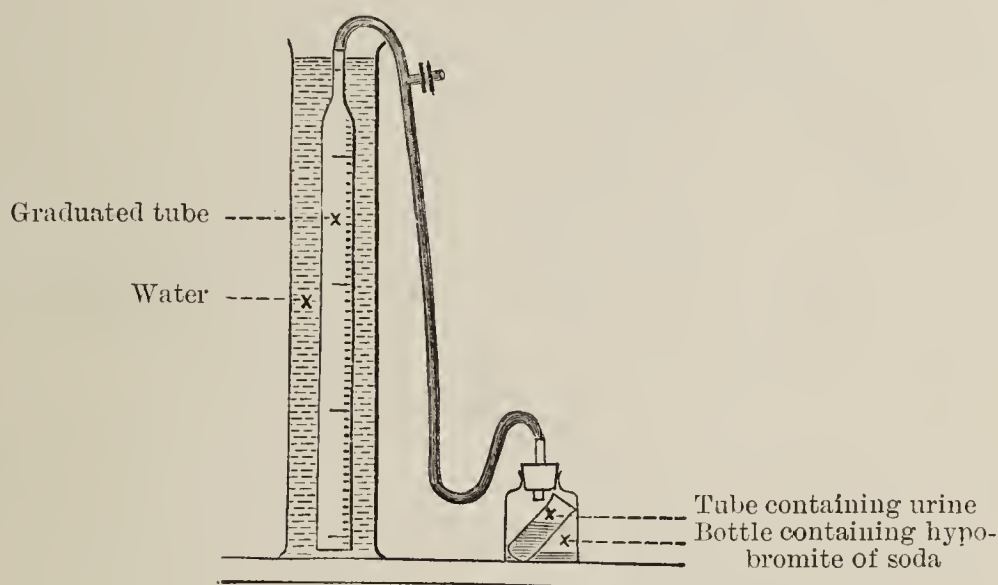


FIG. 101.—ESTIMATION OF UREA BY HYPOBROMITE.

The quantity of urea in a given amount of urine is calculated from the volume of nitrogen evolved when it is decomposed by sodium hypobromite; the apparatus is adjusted so that the two fluids do not mix until required; mixture is effected by tilting the bottle, and the volume of nitrogen is read by means of the graduated tube.

lected during 24 hours, so that the total discharge of urea from day to day can be calculated. The hypobromite method is easy and expeditious, and therefore suitable for clinical use (fig. 101). For more exact determinations, the distillation method in one or other of its numerous modifications must be adopted: it is more laborious, but of more general application, and by it the amount of nitrogen present in any nitrogenous substance—solid or liquid—can be determined (fig. 102).

**Uric acid**,  $C_5H_4N_4O_3$ , which is present to a very small amount in urine (the total excretion not exceeding  $\frac{3}{4}$  gramme per diem), is from a pathological standpoint its most important constituent. An excess of uric acid in the system is the chief physical token of gout, and the



common material of 'gravel,' and of 'calculus,' and of 'chalk-stones.' Normally, uric acid exists in the urine in combination with potassium and sodium as urates of these metals, and, unless the urine be of excessive acidity, it does not crystallise out. But in an unusually acid urine, or in a urine that has been acidified, crystals of uric acid form after a time, which from their appearance have been compared with cayenne pepper; they are diamond-shaped, and their colour is due to the urinary pigments.

Uric acid is insoluble in alcohol, and scarcely soluble in water or in acid solutions, readily soluble in alkaline solutions. The first indication to be fulfilled in cases of gout or gravel is, therefore, to render the urine alkaline. In birds and snakes, uric acid is the chief vehicle of nitrogenous excretion; the excrement of snakes, for example, being little else but uric acid, and having on that account considerable

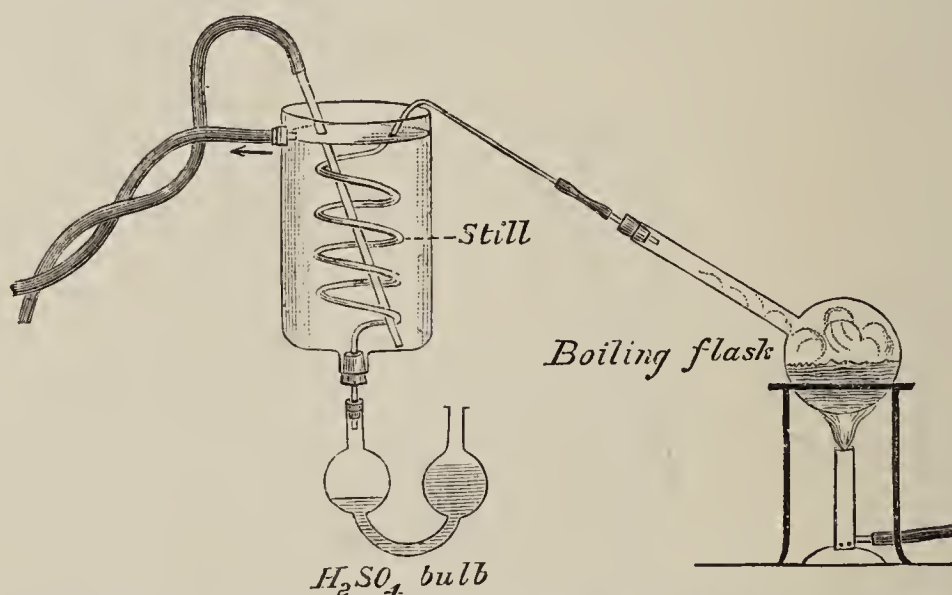


FIG. 102.—ESTIMATION OF NITROGEN BY DISTILLATION OF AMMONIA INTO SULPHURIC ACID. KJELDAHL METHOD; WILFARTH MODIFICATION. (Argutinsky.)

The proteid is oxidised by heating with strong  $\text{H}_2\text{SO}_4$ ; ammonium salts are thus formed; the fluid is then rendered alkaline and heated, ammonia is driven off, and estimated by the loss of acidity in a previously titrated solution of  $\text{H}_2\text{SO}_4$ .

commercial value as a source of uric acid. In mammalia uric acid is a variable product, in herbivora it is replaced by hippuric acid, in the dog (sometimes) by cynurenic acid. Uric acid is a less oxidised product than urea, as may be recognised in the empirical formulæ of the two bodies—urea,  $\text{CON}_2\text{H}_4$ , uric acid,  $\text{C}_5\text{O}_3\text{N}_4\text{H}_4$ ; its physiological antecedents are the still less oxidised bodies, xanthine,  $\text{C}_5\text{H}_4\text{N}_4\text{O}_2$ , and hypoxanthine,  $\text{C}_5\text{H}_4\text{N}_4\text{O}$ ; it is easily oxidised, and can therefore act as a deoxidising or reducing agent, *e.g.* on silver nitrate and on copper sulphate. The action on silver nitrate is utilised as a rough test—uric acid solution dropped on a filter paper, moistened with  $\text{AgNO}_3$ , gives a black or brown stain of reduced silver (Schiff). The action on copper sulphate should be remembered as a possible source of confusion between sugar and uric acid.

*Preparation of uric acid from serpent's excrement.*—Dissolve the

excrement in weak  $\text{NaHO}$  by the aid of heat. Precipitate uric acid from the hot solution with hot  $\text{HCl}$ . Wash with water and dry.

*Detection of uric acid* in the blood and in pathological fluids.—A few drops of blood-serum or other fluid, acidulated, and allowed to evaporate in a watch-glass, in which a linen fibre is laid, deposit crystals of uric acid on the fibre, recognisable by the microscope. This test is applicable to fluids taken from gouty subjects, and is a convenient method for demonstrating the presence of uric acid in the urine.

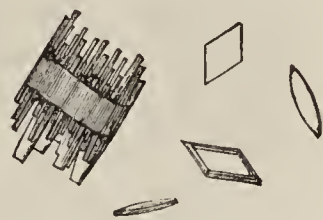


FIG. 103.—URIC ACID CRYSTALS.

*Murexide test*.—Uric acid cautiously evaporated with  $\text{HNO}_3$  gives a yellow stain, which turns purple with  $\text{NH}_3$ , violet with  $\text{KHO}$ .

*Quantitation of uric acid in the urine*.—The clinical significance of uric acid is such as to render an accurate and expeditious method most desirable. No such method exists. After the trial of several, we shall revert to that of Heintz, which, although not to be relied on, is simple and expeditious, and has usually been adopted for clinical purposes. It is as follows:—100 c.c. of urine are acidified by 5 c.c.  $\text{HCl}$ , and set aside for twenty-four to forty-eight hours; the crystalline deposit of uric acid is collected on a weighed filter, washed, dried, and weighed.

**Hippuric acid**,  $\text{C}_9\text{H}_9\text{NO}_3$ , as indicated by its empirical formula, is still less oxidised than uric acid; it is present in considerable quantity in the urine of cattle, is also present in normal human urine, and may be temporarily increased by the administration of benzoic acid. Benzoic acid and glycin form glycobenzoic or hippuric acid, as cholalic acid and glycin form glychocholic acid. Hippuric acid is a crystalline body, soluble in alcohol, and hardly soluble in water; heated in a dry tube it sublimes and deposits benzoic acid, which is also recognisable by its characteristic smell.

*Preparation of hippuric acid from cow's or horse's urine*.—Mix with milk of lime, filter, evaporate—add  $\text{HCl}$ —wash and dry the crystals—redissolve in boiling water—recrystallise.

*From human urine*.—Swallow 1 gramme benzoic acid in a gelatin capsule—collect the urine of the succeeding four hours—proceed as above.

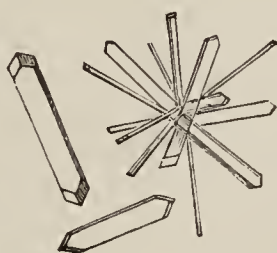


FIG. 104.—HIPPURIC ACID. (From Funke's Atlas.)

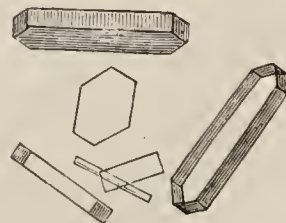


FIG. 105.—CREATIN. (From Funke's Atlas.)

**Creatinin**,  $\text{C}_4\text{OH}_7\text{N}_3$ , present in the urine, is derived from creatin present in muscle and in blood; it is a still less oxidised

product than either uric or hippuric acids. Its presence is recognisable by the following test:—Add sodium nitro-prusside, heat and add  $\text{NaHO}$ ; the solution is ruby-red, soon becoming yellow, and turning green or blue on addition of acetic acid.



*Quantitation of creatinin.* (Neubauer.)—250 c.c. of urine are treated with milk of lime until the reaction is alkaline, then with  $\text{CaCl}_2$  until no further precipitate appears. The filtrate is evaporated on a water-

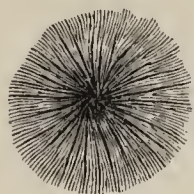


FIG. 106—CREA-  
TININ - ZINC-  
CHLORIDE.

bath to a syrupy consistence, and extracted for several hours with strong alcohol. To the alcoholic extract are added a few drops of a neutral alcoholic solution of  $\text{ZnCl}_2$ , and the beaker is set aside for three days in a cool place. A crystalline precipitate of creatinin-zinc-chloride, in the form of rosettes, visible to the naked eye, is found at the bottom and sides of the beaker.

This precipitate is collected on a weighed filter, which is subsequently dried and weighed.

The bodies above considered—urea, uric acid, hippuric acid, and creatinin—are the chief nitrogenous derivatives present in normal urine; urea is the most fully oxidised, creatinin the least fully oxidised; uric acid occupies an intermediate position. A further indication of the relationship between these several bodies is furnished by the fact that urea can be obtained from uric acid by oxidation, and from creatin by treatment with alkalis.

**Inorganic salts.**—The chief inorganic salts of the urine are *chlorides*, *phosphates*, and *sulphates*. Of these, the sulphates are of most physiological interest, for they represent the form in which nine-tenths of the sulphur contained in proteid food escapes from the body; they are not simply reappearing sulphates swallowed with food, for the amount of food sulphates is so small as to be negligible. Chlorides and phosphates, on the other hand, proceed from chlorides and phosphates swallowed with the food. The notion that the amount of phosphate excreted varies with the degree of mental exertion is entirely unsupported by facts, and when we realise that the amount of phosphoric acid in the whole cerebro-spinal substance is only  $\frac{1}{10}$  the amount in the muscles, and  $\frac{1}{100}$  the amount in the bones, there is no ground for expecting to find variations of nerve metabolism represented by variations of phosphoric acid in the urine. The average daily excretion of the three acids in the form of salts is  $\text{HCl}$ , 7.5 grms.;  $\text{H}_3\text{PO}_4$ , 3 grms.;  $\text{H}_2\text{SO}_4$ , 3 grms. The excretion of chlorides is said to be diminished in pneumonia; that of phosphates is said to be increased in fever and in bone disease, diminished during pregnancy.

**Chlorides** (Na and K).—The presence of chlorides is recognised by the precipitate formed with silver nitrate, insoluble in nitric acid.

*Quantitation* (Mohr).—The principle of the method is as follows:—Chlorides are precipitated by a standard solution of silver nitrate (29.063 grms. in 1,000), of which 1 c.c. will precipitate .01 gm.  $\text{NaCl}$ ; potassium chromate is used to indicate when an excess of silver has been reached. Take 10 c.c. of urine and dilute ten times with distilled water, add a few drops of potassium chromate solution; run the silver solution into this mixture until an orange colour appears, showing that

the excess point has been passed. The method is expeditious, and therefore suitable for clinical work, but it is not very exact; the readings are generally 1 c.c. too high, and should be corrected by that amount. If, for instance, the excess point is passed at 6 c.c. of silver solution, the 10 c.c. of urine are calculated as containing .05 gm. NaCl, or .5 per 100.

**Phosphates** (Mg, Ca, Na, K).—The addition of ammonia to urine gives a precipitate of earthy phosphates (Ca and Mg); the filtrate from this, or the original urine, gives a precipitate of alkaline phosphates (Na and K) on boiling with ammonium molybdate and nitric acid.

The amount of phosphoric acid present is determined as follows:—

Two solutions are prepared. (1) *Standard uranium acetate solution*, 35 gm. to 1,000 c.c. water with 25 c.c. glacial acetic acid. 1 c.c. of this solution is nearly equivalent to .005 gm.  $P_2O_5$ . (2) A fresh 10 per 100 solution of potassium ferrocyanide to be used as the *indicator*.

Heat 50 c.c. of urine with successive additions of the standard solution from a graduated pipette, until a drop of the mixture strikes brown with a drop of the indicator; *e.g.* if the brown colour is obtained after 20 c.c. of the standard solution have been added, the 50 c.c. urine contain .100 gm.  $P_2O_5$ , *i.e.* .2 per 100. The uranium solution should be previously adjusted so that 20 c.c. will exactly balance 50 c.c. of a .2 per cent.  $P_2O_5$  solution (ammonio-sodic phosphate 5.886 gm. in 1,000 water).

The principle upon which the determination depends is that uranium nitrate combines with phosphoric acid in a definite ratio, the point of excess being indicated by the brown colour which *free* uranium nitrate gives with potassium ferrocyanide.

**Sulphates**.—Sulphur (mainly derived from proteid) escapes in the urine in the form of sulphates:—(1) In chief part as inorganic sulphates (of K and Na). (2) In small part as aromatic sulphates, *viz.* an indigo-yielding salt, potassium indoxyl sulphate,  $C_8H_6NKSO_4$ ; and a phenol-yielding salt, potassium phenyl sulphate,  $C_6H_5KSO_4$ .

Under normal circumstances the amount of aromatic sulphates present in the urine is about  $\frac{1}{10}$  that of the inorganic sulphates, but abnormally, when putrefaction of proteid occurs in the small intestine, leading to the formation of aromatic bodies such as indol and phenol, this proportion may be considerably increased. The total amount of sulphates depends upon the amount of proteid consumed; the average daily excretion of 3 grms.  $H_2SO_4$  contains about 1 gm. S., *i.e.* about as much as is contained in 100 grms. of proteid.

The *presence* of sulphates is recognised by the precipitate formed with barium chloride, insoluble in hydrochloric acid. The indigo-yielding substance (so-called 'indican') is present in excess if the urine on boiling with HCl gives a pink colour. Its physiological antecedent is indol, formed in the intestine by the putrefaction of



proteids. To demonstrate its presence in ordinary urine, a considerable quantity, *e.g.* 500 c.c., should be taken, and to it 250 c.c. pure HCl be added. A coppery scum forms on the surface. The greater part of the fluid is syphoned off, and the remainder, including the scum, is thrown on a filter. The filter is washed with ammonia, and then with alcohol. The alcohol takes up indigo and becomes red; on boiling it becomes blue.

To estimate the *total amount* of sulphates (inorganic and aromatic), add 5 c.c. HCl to 100 c.c. urine, and heat in order to break up aromatic sulphates; precipitate with  $\text{BaCl}_2$ , and heat on water-bath; filter, wash with water and with alcohol, dry and weigh. Supposing the filter to have gained .5 gm. in weight by the  $\text{BaSO}_4$  precipitate, we ascertain by calculation that the amount of  $\text{H}_2\text{SO}_4$  present is .21 gm. and of S about .07 gm. To estimate the amount of aromatic sulphates alone, the same process is gone through with the *filtrate* of urine from which the inorganic sulphates have been removed by baryta mixture ( $\text{BaO}$  and  $\text{BaCl}_2$ ), without previous heating with HCl. The weight of the  $\text{BaSO}_4$  precipitate is more exactly obtained by igniting the filter paper and precipitate in a covered platinum capsule, and after cooling in an exsiccator, weighing the ash in the capsule; the weight of the capsule and that of the filter-paper ash are known, the remainder is the weight of  $\text{BaSO}_4$  (Salkowski).

A more expeditious method is by volumetric determination of the amount of  $\text{BaCl}_2$  required to precipitate all the sulphates present. Each c.c. of the standard barium solution (30.5 gm.  $\text{BaCl}_2$  in 1,000) will precipitate .01 gm.  $\text{SO}_3$  as  $\text{BaSO}_4$ . The excess point is indicated by the aid of a 20 per cent. solution of potassium sulphate; a drop of this solution mixed with a drop of the clear stratum of the urine under treatment should give no more than a slight cloudiness; if a clear drop of the latter, mixed with a drop of  $\text{BaCl}_2$  solution, gives a precipitate, the process is not yet complete; if it becomes milky with a drop of the indicator, too much  $\text{BaCl}_2$  has been added. The urine is to be heated before each addition of  $\text{BaCl}_2$ . If it has been previously boiled with HCl, the total sulphates are determined; if not so treated, only the inorganic sulphates. Supposing the excess point to be obtained with 15 c.c. of barium solution added to 100 c.c. urine, the percentage of  $\text{SO}_3$  is known to be .15 (= nearly .2  $\text{H}_2\text{SO}_4$ ).

**Abnormal constituents.**—The substances that most commonly appear in the urine under abnormal circumstances are *albumin*, *blood*, and *sugar*; and the part of the kidney by which they are believed to escape is the glomerulus. In jaundice the urine is tinged with *bile-pigment*.

**Albumin** may leak from the blood into the urine under several conditions and in several forms. It may appear as serum-albumin, serum-globulin, albumose, or peptone. Its presence may be due to serious structural alteration of the kidney, as in Bright's disease, or

to a trifling and evanescent alteration of function, as by diet, or in a condition of fatigue, and experimentally it may be induced by altering the state of the blood, or by interfering with the circulation of the kidney. The readiest and most certain test for albumin in the urine is heat *plus* nitric or acetic acid. If the acidulated urine becomes turbid, or coagulates on boiling, it contains albumin. The degree of the turbidity or the amount of the coagulation gives a measure, though a very imperfect one, of the amount of albumin. To make a correct estimate the acidified boiled urine should be thrown on a weighed filter, which is subsequently dried and weighed.

The non-escape of albumin under normal conditions is to be ascribed to the integrity of the glomerular epithelium, and to a normal state of the blood. Interference with the nutrition of the kidney, as by temporary arrest of its circulation, alteration of the composition of the blood, as by injection of water or of lake blood, are apt to provoke albuminuria or even hæmoglobinuria.

**Sugar.**—The *presence* of sugar is readily ascertained by *Trommer's test*. A urine that contains sugar gives a red or yellow precipitate on boiling with caustic potash and a few drops of copper sulphate; a preferable mode of testing is to add urine to Fehling's fluid that has just been boiled, heating the mixture again to near boiling-point.

*Fischer's test* is a valuable means of identifying sugar in the urine, inasmuch as it yields a negative result with other reducing substances, such as creatinin and uric acid. A little powdered *phenylhydrazin-hydrochloride* and sodium acetate are added to the suspected fluid, which is boiled and left for half an hour in a hot-water bath; a crystalline deposit is formed, microscopically, as bundles of yellowish needles. Glycuronic acid gives the same result, but does not ferment with yeast.

Barfoed's reagent (neutral acetate of copper, 6·6 grms. ; acetic acid, 2·5 c.c. ; water, 100 c.c.) serves to distinguish dextrose from other reducing sugars (lactose, maltose, &c.) ; dextrose alone gradually effects reduction of this fluid, without boiling.

The *amount* of sugar present may be determined in several ways, of which the following are the principal :—(1) by *Fehling's solution*, (2) by *fermentation*, (3) by the *polarimeter*. Of these methods, the most exact as well as the most delicate is the first, which is as follows :—Fehling's solution<sup>1</sup> is prepared of such strength that the copper salt in 10 c.c. is reducible by  $\frac{1}{20}$  gm. sugar ; the completion of reduction is indicated by disappearance of the blue colour of the solution. A measured quantity of Fehling's solution is boiled with urine, which is added until the blue colour disappears. For clinical use the determination can be conveniently carried out in a test-tube with 1 c.c.

<sup>1</sup> To make *Fehling's solution*.—(1) Dissolve 34·64 grms. pure dry powdered  $\text{CuSO}_4$  in 200 c.c. distilled water. (2) Dissolve 173 grms. Rochelle salt (sodio-potassic tartrate) in 480 c.c.  $\text{NaHO}$ , S.G. 1·14. Mix and dilute to 1,000 c.c.



of Fehling, and it will usually be necessary to dilute the urine; for more exact determinations 10 c.c. should be employed, and the boiling carried on in a capsule into which the urine is dropped from a burette. If, for instance, 1 c.c. is decolorised by 4 c.c. of a urine that has been diluted ten times, the original urine is known to contain 1·25 per cent. of sugar; if the day's urine amounted to 2,000 c.c., then the sugar passed in that day would amount to 25 grammes. In extreme cases of diabetes much higher amounts have been passed per diem, up to 500 or even 750 grammes, *i.e.* more than a pound.

*Fermentation test.*—The sugar in a diabetic urine undergoes fermentation if left in a warm place mixed with a little yeast; the sugar is decomposed into carbon dioxide and alcohol, and the specific gravity of the urine is lowered. The amount of sugar can then be roughly gauged; each degree of S.G. lost represents about 1 grain of sugar to the ounce of urine; *e.g.* a loss of 5° S.G. and a daily discharge of 80 ounces of urine gives the day's loss of sugar as 600 grains = about 26 grammes.

The *polarimeter* is applicable only to very strongly saccharine urine, which should be previously decolorised. Diabetic sugar is dextro-rotatory, and a column of fluid 20 cm. long will rotate the plane of polarisation about 1 degree for each gramme of sugar per cent.

The tests by which blood- and bile-pigments are recognised have been given at pages 39 and 200.

**Micturition** is a reflex act that can be permitted or resisted by voluntary action. The normal excitant of the act is the distension of the urinary bladder, causing, in consciousness, a desire to micturate. The voluntary response to this sensation may be 'yes' or 'no'—'yes' by contraction of the abdominal muscles, with a passive condition of the sphincter, 'no' by contraction of the sphincter, with a passive condition of the abdominal muscles. In the latter case the desire is resisted, and relief is deferred. In the former case the desire is satisfied by a normal act of relief, which commences with contraction of the bladder and of the abdominal muscles, and ends by contraction of the sphincter vesicæ and of the urethral muscles (bulbo-cavernosus—muscular coat of urethra). In its performance an excitation of the vesical end of the urethra by the first drops of urine acts as a further stimulus to the reflex act, which may still be inhibited, although with more difficulty.

It may happen—

1. That the excitation is excessive, the desire imperative, the voluntary negative insufficient to arrest the act. Involuntary micturition then occurs, particularly in children.

2. That the sphincter is paralysed. Incontinence of urine is the effect.

3. That the bladder, or that the abdominal muscles, or that both, are paralysed; retention of urine in a distended bladder, followed by overflow (= incontinence), is then produced.

4. That the muscles act in an inco-ordinate manner. This condition, which is analogous with 'stammering,' causes temporary retention. Like stammering, it is apt to be produced by emotion.

The mechanism of micturition is thus in most respects similar to that of defæcation (*vide* p. 169), the particular points of difference being referable to the fact that whereas the large intestine is of such great capacity that it can retain the fæcal accumulation of many days, or even weeks, the bladder is over-filled by the urine secreted in a single day. Consequently, the two opposite conditions, retention and incontinence, are usually associated; retention leading to overflow or incontinence, incontinence being the effect of retention. Moreover, micturition is more subject to voluntary control and less influenced by habit than is the case with defæcation.

*Vesical pressure.*—The passive pressure of fluid in a full bladder amounts to about 1 cm. Hg, and it has been found that the sphincter at rest, by virtue of its elasticity alone, can withstand a vesical pressure of about 3 cm. Hg. A normal man, by the effort of micturition in which the bladder and abdominal muscles are contracted, can raise the pressure to about 10 or 12 cm. A paraplegic patient by the action of the bladder alone can raise it to 4 or 5 cm. The vesical pressures just referred to are not to be confused with the secretory pressure of the kidneys, which, as measured by a manometer connected with a divided ureter of the dog, has been found to reach to about 6 cm.

**Excretory action of the liver.**—Allusion has been made to the part that the liver plays in excretion; it was mentioned that the bile is not only a secretion serving in digestion, but to some extent an excretion to be got rid of, the most distinct proof of the fact being the identity of urobilin with hydrobilirubin; it was mentioned, moreover, that the liver forms not only bile-products (bile-pigments and bile-acids), which are carried away in the bile, but also urinary products (urea, hippuric acid, uric acid), which are carried away in the blood, and separated by the kidneys. We have now to examine this physiological association between liver and kidney as excretory organs—an associa-



tion that we may at once characterise as formation by the liver, separation by the kidney. The chief items in the argument are the following:—1. Bilirubin is made by the liver ; urobilin is discharged by the kidney. 2. Urea is made by the liver ; it is discharged by the kidney. 3. Glycin is made by the liver ; hippuric acid (which is glycin + benzoic acid) is discharged by the kidney. 4. Uric acid is made by the liver, discharged by the kidney. 5. Sugar, made by the liver, may be discharged by the kidney.

The data upon which these several dicta are based are as follows:—1. Urobilin is chemically identical with reduced bilirubin, or hydrobilirubin. 2. A freshly-excised liver, submitted to artificial circulation with blood containing ammonium carbonate, yields urea found in the outcoming blood. In acute atrophy of the liver, with abolition of hepatic function as its consequence, leucin and tyrosin are found in the urine in place of urea. 3. Benzoic acid taken by the mouth appears as hippuric acid in the urine, but after excision of the liver it passes unchanged as benzoic acid. 4. After excision of the liver on geese, uric acid ceases to be excreted (Minkowski). 5. Excessive sugar-formation by the liver is accompanied by discharge of sugar by the kidney.

We must not, however, take the association between liver and kidney—viz. formation by the former, separation by the latter—in too absolute a sense, as implying a strict division of labour. Urea and hippuric acid are not formed by the liver exclusively ; other organs—the kidney in particular—submitted to artificial circulation, are capable of forming these substances, this being especially the case as regards the formation of hippuric acid by the kidney. Blood containing benzoic acid and glycin, made to pass through a recently excised kidney, or left in contact with a minced fresh kidney, is after a few hours found to contain hippuric acid. Thus the living kidney can effect the conjugation of glycin (formed by the liver) with benzoic acid (formed or set free in the course of digestion). The chief evident function of the kidney is the separation of formed materials, but it certainly also contributes to the actual formation of the materials that it separates.

On the other hand, we must recognise that the liver, although it is pre-eminently formative of excreta, such as urea and (in birds) uric acid, is not thus formative to the entire exclusion of other organs and tissues. We have already seen an illustration of this in the case of bile-pigment ; although no doubt the principal normal relation is formation of bilirubin by the liver,

and separation of urobilin by the kidney, yet we must admit that, failing the liver, urobilin may take origin otherwise in the body.

**Diabetes.**—The essential phenomena of diabetes are the passage of sugar into the urine, and an excessive discharge of water. The first event is the characteristic of *diabetes mellitus*, and is found to be associated with the presence of an excessive amount of sugar in the blood; the second event is the characteristic of *diabetes insipidus*. No conclusive proof has yet been given to show that the urine normally contains any sugar, and it is certain that normal urine does not contain it in considerable amount; in a case of *diabetes mellitus* the urine may contain as much as 5 per cent. The average percentage of sugar in normal blood is only  $\cdot 1$ ; the percentage of sugar in diabetic blood may reach  $\cdot 5$ . Comparing these two amounts,  $\cdot 5$  per 100 in blood, 5 per 100 in urine, we recognise that mere diffusion will not account for the excretion of sugar, but that, as in the case of urea, an activity of renal epithelium must be invoked. The excess of sugar in the blood, disposed of by the kidney, is attributable to two causes:—(1) abnormal action of the liver, (2) deficient consumption by the tissues; and it is probable that both causes contribute to the result. The abnormal action of the liver can be regarded in two different lights: either the liver action is defective, sugar coming from the intestine is not stored as glycogen; or the liver action is excessive, more sugar than usual is made. The second alternative is the more probable; diabetes is not produced by diseases in which liver action is certainly defective, and it may persist, though in diminished degree, when all starch and sugar are withheld from the dietary.

The relation between excessive action of the liver and diabetes is further borne out by the effects that follow puncture of the spinal bulb, and several other operations. Rabbits shortly after a puncture of the fourth ventricle become diabetic in so far as they pass sugar in the urine. Frogs similarly treated, especially at the beginning of winter, are similarly affected. This diabetic state is, however, temporary, and an essential condition of its appearance is that the liver shall, at the time of puncture, contain a large store of glycogen. It appears, therefore, that *puncture diabetes* is due to an accelerated conversion of liver-glycogen into sugar. This view is borne out by further observations; it has been found that several other operations will produce the same result, *e.g.* section of the cervical sympathetic or of the spinal cord, destruction of the



cervical or of the stellate ganglia, excitation of the central end of the vagus, or indeed of any afferent nerve, ether, morphia, apnœa, and, finally, the injection of arterial blood into the portal vein; the common effect in all these cases, viz. sugar in the urine, being attributable to one common cause, viz. to hyperæmia of the liver. It should be added, however, that the sugar reaction may be simulated by the allied carbohydrate, glycuronic acid,  $C_6H_{10}O_7$ , which, according to Ashdown, is the substance giving the apparent sugar reaction in the urine of chloroform narcosis. Puncture of the bulb may also give rise to *diabetes insipidus*; this effect, the mechanism of which is more obscure, is attributable to vasomotor disturbance of the kidney rather than of the liver.

*Pancreatic diabetes.*—Recent experiments have shown that the pancreas bears some peculiar relation to the sugar function; dogs are rendered diabetic by complete removal of the pancreas, the diabetes so produced remaining permanent until death (Minkowski and v. Mering), contrasting in this respect with the temporary character of *puncture diabetes*, as well as in the fact that it is not a nerve effect; the nature of the relation between diabetes and loss of the pancreas is as yet entirely obscure; it is comparable with the equally obscure action of the thyroid gland, *i.e.* in both cases removal of a gland causes a peculiar malnutrition, characterised by excess of sugar if the pancreas is removed, by excess of a mucinoid substance (myxœdema) if the thyroid is removed; in both cases, therefore, the gland in question must be of importance to healthy nutrition, possibly by virtue of some as yet unknown process of ‘internal secretion.’ This is not the place to discuss at length the possibilities that are suggested by observations of this order; it is enough for us to recognise that a relation—however obscure—really subsists between the special functions of particular organs and the general nutrition of the body, and that remedial measures may be looked for by following up this clue; successful results are, in fact, reported subsequent to injections of testicular, pancreatic, and thyroid extracts in cases where the function of the respective glands was presumably deficient; experimentally, dogs rendered diabetic by removal of the pancreas have had life prolonged by injection of pancreatic extract, and by pancreatic grafting.

These are modern examples of the conception long ago formed by Treviranus, to the effect that the several parts of the body are mutually interdependent, the waste of one organ serving as the raw material to another organ, and even the organs themselves being conceivably ‘excretory products’ in relation to the remainder of the organism. The development of the mammary gland coincident with the presence of a foetus, the arrested development of male characteristics in consequence

of castration, may also be pointed to in token of the obscure inter-organic relations which play a part in the internal economy of the processes of nutrition.

‘Addison’s disease,’ of which the most obvious sign is a bronzed skin due to excessive pigmentation, is perhaps another case in point; the pigmentation is one among other signs of a malnutrition which, in some cases at least, is attributable to diseased suprarenal glands. Schäfer and Oliver have shown that an injection of suprarenal extract (of the medulla, not of the cortex) causes a marked increase of blood-pressure.

Even the kidney, which is principally concerned with a terminal act of metabolism, is perhaps to be recognised as playing a part in its earlier stages. According to Bradford the disturbance of nutrition caused by partial removal of the kidneys is altogether different from that caused by their entire removal. His observations are to the following effect:—A large wedge of one kidney, amounting to between  $\frac{1}{2}$  and  $\frac{2}{3}$  its total weight, is excised, the surfaces of the remaining portion are fastened together; when the animal (dog) has recovered, the entire second kidney is removed, thus leaving only  $\frac{1}{3}$  to  $\frac{1}{2}$  a single kidney on duty; the consequences of this changed condition are: *acute emaciation* (e.g. a fall of body-weight amounting to 37 per cent. in fourteen days), in spite of normal appetite and diet; *greatly increased discharge of water*, and, in less marked degree, *increased discharge of urea*. These effects seem to suggest that the renal function is not merely to remove degradation products, but also to produce some as yet unknown body or bodies, adjuvant of tissue-integration, or obstructive of tissue-disintegration. We must recognise that our rough fragments of precise knowledge, in terms of chemistry and physics, cover but a small portion of the phenomena actually accomplished within the living body. And we should above all keep our minds open to the idea that the several organs of the organism are interdependent—many members of one body.

**Excretory action of the skin.** SWEAT.—The *skin*, by virtue of its glands, is an excretory organ. The substance excreted is almost exclusively water, the amount discharged varying to such an extent with temperature as to cause the urine to be pale and copious in winter, dark and scanty in summer. Sweat, when perfectly fresh, is alkaline, but this reaction is very quickly reversed owing to fatty acids produced in its decomposition.

The connection between the cutaneous and renal channels of excretion is further illustrated by the appearance of urea in the sweat; even normal sweat is said to contain a trace of urea, and the sweat of a uræmic patient may contain the substance in considerable quantity.

The amount of evaporation taking place from the surface of



the body depends upon the physiological activity of the sweat-glands, upon the movements of respiration, and upon the physical state of the atmosphere as regards temperature and pressure, saturation with moisture, and movement of the air. Low temperature, saturation, and stillness are unfavourable, the reverse conditions are favourable to evaporation. Under ordinary conditions air contains about 1 vol. per cent. of water vapour. Air at normal pressure is *saturated*

at	0°	by	·65	per cent.	water vapour.
„	10°	„	1·3	„	„
„	20°	„	2·3	„	„
„	30°	„	4·15	„	„
„	40°	„	7·25	„	„

*e.g.* we have seen that the expired air is saturated at 37°, *i.e.* it contains about 7 per cent. water vapour; if into an atmosphere at 1 per cent. saturation a man expires per diem 10,000 litres air, he gives off 600 litres water vapour (= 480 grammes).

From the skin the water is given off partly by ‘insensible perspiration,’ partly as visible drops of sweat, and the latter partly evaporates from the surface, partly trickles off or is absorbed by clothing. The daily amount of perspiration may be put at 900 litres of water vapour (= 720 grammes). Its rate of discharge in the form of insensible perspiration may be gauged by a hygrometer, consisting of a shallow glass vessel coated with calcium chloride, which is weighed before and after it has been left inverted over the skin for a given period; the increase of weight indicates the amount of water vapour absorbed by the calcium chloride (w.). Tested in this way the palm of the hand is found to give off moisture more rapidly than any other equal area of skin. The constant transpiration of water from the skin and lungs, and the greater *weight* of CO<sub>2</sub> output as compared with O<sub>2</sub> income, cause an appreciable fall of body-weight, *i.e.* nearly 1 gramme per minute, or 2 ounces per hour.

The secretion of sweat is under the control of sympathetic nerve-fibres which leave the spinal cord by the anterior roots and run a similar course to that of vasomotor nerves. If the peripheral end of the sciatic nerve of a young cat be excited, beads of sweat break out from the pad, and the effect can still be obtained after arrest of the circulation or after amputation of the limb; thus proving the separate existence of sudo-motor nerves (Luchsinger). As in the case of the salivary gland, pilocarpine

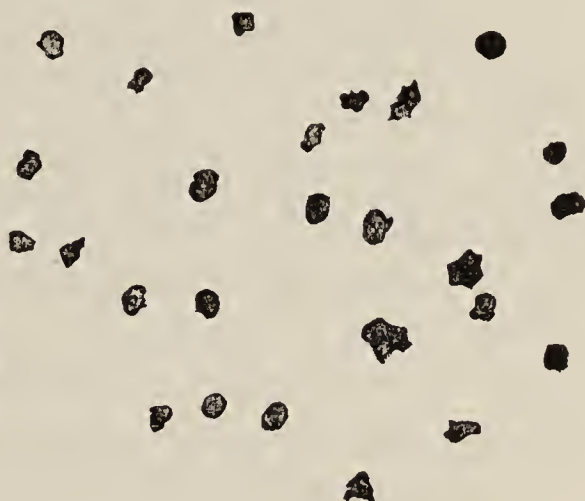
stimulates the secretion and atropine stops it. Langley and Sherrington have recently shown that the sympathetic also has action upon the muscles of the hair roots ; on the cat stimulation of the lowest sacral and coccygeal roots, or ganglia, causes the hairs of the tail to bristle up to a very unmistakable degree.

An elegant demonstration of the local action of atropine upon the sweat-glands has been given by Aubert :—A sheet of white paper, previously dipped in a .25 per cent. solution of silver nitrate, applied to the skin for a minute or two and then exposed to the light exhibits black dots or ridges wherever the nitrate has been acted upon by the chlorides contained in the sweat issuing from the glands ; if this be done upon a portion of skin including a patch upon which belladonna plaster (atropine is an alkaloid derived from *atropa belladonna*) has been left for twenty-four hours, a blank patch will appear in the sweat-print, corresponding with the atropinised spot.

We may take this opportunity of mentioning the changes of hue that the skin undergoes in many animals according to



PIGMENT CELLS DIFFUSE.  
In darkness.—Moderately dark skin.



PIGMENT CELLS CONCENTRATED.  
In daylight.—Moderately light skin.

FIG. 107.

Low-power view of Frog's skin to illustrate the effect of light upon the pigment-cells. (Lister.)

the light. A common frog (*R. temporaria*) kept in the dark has a dark skin ; if exposed to light the skin soon becomes pale ; the change depends upon the greater or smaller diffusion of pigment granules within branched pigment-cells (chromatophores), the pigment being widely diffused with dark skin, collected together in small lumps in a pale skin. The pale state is thus analogous to the contracted state of muscle ; it may be produced by arrested circulation, or at death, or by excitation of the sciatic nerve, and apparently the effect of light is partly direct, partly reflex. (See also Dubois' observations, p. 456.)



## CHAPTER VII

## FOOD, NUTRITION, AND EXCRETION. ANIMAL HEAT

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## ANIMAL HEAT

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THE living body yields *energy* in the form of *work* and of *heat*. This result is brought about by chemical action. Matter is used up and becomes useless, organic compounds disintegrate and their waste products are got rid of. Consumption of matter is thus a primary condition of vital activity, and it is necessary to the continuance of life that new matter should take the place of

spent matter. The food—proteids, fats, and carbohydrates—supplies the necessary new matter, and is the primary source of all body energy, by its transformation into work, heat and waste products. The essential waste products are *urea* and *carbon dioxide*; the essential elements in these waste products as well as in the food are *nitrogen* and *carbon*.

**From proteid to urea.**—*Nitrogen* enters the body in *proteid*, and leaves it in *urea*. In what state or states does it exist in the body, how does nitrogen traverse the body? The answer is necessarily fragmentary and in many respects hypothetical. Proteid by digestion becomes peptone, which is absorbed and retransformed into the proteids of plasma and of lymph. These fluids surround and permeate the organised elements of the tissues, and the proteid that they carry forms the floating balance of nutritive matter from which the comparatively fixed capital of living protoplasm is supplied. Of this *free, circulating or coasting proteid* in blood and in lymph, it is supposed that only a small proportion is actually taken into chemical combination in protoplasm as fixed or *organ proteid*, and that the greater proportion is acted upon and used by living protoplasm without being integrated by it to make part of its own substance. The free or coasting proteid thus used up gives rise to urea, as does that small proportion of fixed or organ proteid which disintegrates and gives place to a correspondingly small proportion of newly-integrated proteid. It is supposed that if a superfluous amount of proteid be swallowed, the excess may in the intestine itself, by the agency of pancreatic juice, be carried beyond the peptone stage and become leucin and tyrosin, which, being absorbed into the portal system, may be at once converted into urea by the agency of the liver. Thus, from proteid to urea there are three usual roads: (1) the short-cut *via* leucin and tyrosin in the intestine, and urea in the liver; (2) the high-road *via* circulating proteid; (3) the long, narrow way *via* circulating and organ proteid. The centre of action is the living tissue-element, which, while undergoing little change as to its own proteid, effects considerable change of the proteid solution that soaks through and around it. An analogy may serve to illustrate these alternatives. A sovereign may (1) be changed and spent at once (= excess proteid), or (2) be sent to the bank to form part of a balance that is being used up as required (= circulating proteid), or (3) be invested in comparatively permanent capital (= organ proteid).



*Intermediate substances between proteid and urea.*—Can we identify any named substances intermediate between proteid and urea? if so, have we any reason to admit that the main current of nitrogen passes in such substances? The second portion of the question may at once be answered in the negative; we find incoming nitrogen in proteid, outgoing nitrogen in urea, but we possess no data enabling us to trace the main current of nitrogen through any named intermediate stage between proteid and urea.

But we may not on this account assert that there are no such intermediate stages, and that urea is immediately derived from proteid; we are to accept the fact that the nitrogenous current is concealed during its passage through the organism, and that we do not know what forms and channels it may take in that passage. It is, however, not totally concealed; we can, among the nitrogenous principles of the body, recognise two or three named bodies in small quantities, less complex than proteid, more complex than urea, and it is allowable to suppose that these bodies are intermediate between proteid and urea, although we are not able to say whether one, or all, or none are, so to speak, on the main line from proteid to urea. Bodies belonging to this category are *leucin* and *tyrosin* (in the intestine), *glycin* (in glycocholic acid), *creatin* and *sarcosin* (in muscle), *uric acid* and *hippuric acid* (in the urine).

The relations of leucin and tyrosin to proteid on the one hand and to urea on the other have been already discussed; these bodies are, properly speaking, formed ‘outside’ the body, although within the digestive tract, and are in no sense to be placed as intermediate stages between proteid in the body and urea emerging from the body. They are named here chiefly because they can appear in the urine in place of urea.

*Glycin* occurs in glycocholic acid, which is secreted by the liver, re-absorbed from the intestine, and excreted in part as hippuric acid, in part as urea.

*Creatin* is a component of muscle; it is chemically convertible into urea: 
$$\text{C}_4\text{H}_9\text{N}_3\text{O}_2 + \text{H}_2\text{O} = \text{CON}_2\text{H}_4 + \text{C}_3\text{H}_7\text{NO}_2,$$
 but no proof has yet been given that the conversion takes place as a physiological event in the body.

*Uric acid* is a less fully oxidised product than urea, and may wholly or partially take its place; the nitrogenous excretion of

birds and of serpents occurs entirely in the form of uric acid; and in the human subject a characteristic feature of gout is an increased formation of uric acid and a deficient discharge of urea. There is evidence to show that the series:—uric acid  $C_5H_4N_4O_3$ , xanthin  $C_5H_4N_4O_2$ , and hypoxanthin  $C_5H_4N_4O$ —arises from the decomposition of nucleo-albumin; but there is no proof that uric acid is a normal antecedent of urea.

**From fat and carbohydrate to carbon dioxide.**—Carbon enters the body in fat and in carbohydrates (starch and sugar), and leaves it in carbon dioxide. It must also be remembered that some carbon enters the body in proteid, and that some hydrogen from fat must leave the body in water. What are the intermediate stages between the initial terms—fat, starch, and sugar—and the final term, carbon dioxide? The answer to this question is necessarily imperfect, for carbon, while in the organism, is hardly less concealed from us than nitrogen. The best-attested probabilities are:

1. Carbon in proteid may become carbon with oxygen ( $CO_2$ ), immediately by the disintegration of proteid, or mediate after the previous formation of fat or of glycogen from proteid.

2. Fat absorbed can exist in the body only as fat, and cannot give rise to proteid or to carbohydrate. The destiny of fat, whether recently absorbed or previously deposited, is to be oxidised, yielding  $CO_2$  and  $H_2O$  as its final products. Absorbed fat may thus be immediately expended by oxidation, or it may be stored in the body as a more or less permanent reserve of fat, which becomes oxidised when required. Fat so stored as adipose tissue is, although 'fixed,' analogous with circulating albumin in this respect, that it is mainly a neighbour and not a constituent of protoplasm, and there is no positive evidence that fat can contribute to the formation of proteid or be converted into carbohydrate.

3. Carbohydrate absorbed (in the form of dextrose or lævulose) may be immediately oxidised, yielding  $CO_2$ , or it may be stored as a comparatively temporary reserve of glycogen in the liver. There is no evidence that carbohydrate forms proteid, and it is not absolutely proved that it is directly converted into fat in the body, though it certainly favours the deposition of fat (*vide* p. 261). In the muscles it is a possible source of lactic acid, which by further oxidation yields  $CO_2$ ; it must be admitted, however, that recent observations do not confirm this view, which is in harmony with the accepted doctrine of carbohydrate consumption



in muscular activity; on the other hand, the alternative supposition that lactic acid proceeds from disintegrated proteid is not definitely proved. The above statements, which are no more than a brief summary of the best authorised facts and opinions concerning the general drift of the animal food-supply, may be still further summarised as follows :—

Food	State in Body	Chief excretion
Proteid . . .	$\left\{ \begin{array}{l} \text{Proteid } \left\{ \begin{array}{l} \text{circulating} \\ \text{organised} \end{array} \right. \\ \text{Glycogen and glucose} \\ \text{Fat} \end{array} \right.$	$\left\{ \begin{array}{l} \text{CON}_2\text{H}_4 \\ \text{CO}_2 \\ \text{CO}_2 \end{array} \right.$
Fat . . .	Fat	CO <sub>2</sub>
Carbohydrate . .	$\left\{ \begin{array}{l} \text{Glycogen and glucose} \\ \text{Fat? Lactic acid?} \end{array} \right.$	$\left\{ \begin{array}{l} \text{CO}_2 \\ \text{CO}_2 \end{array} \right.$

Turning our attention to a collateral aspect of the same sequence, viz. to the elementary composition of proteids, fats, and carbohydrates, we shall more fully appreciate two distinct and important points—(1) that a proteid molecule can conceivably be split into a nitrogenous and a non-nitrogenous moiety, as has just been stated; (2) that the ‘respiratory quotient’ must vary with the nature of the diet, and how so, this being the same problem from its other side as that which has already been presented to us at p. 134.

(1) *The carbon surplus*.—The nitrogen contained in 100 grammes of dry proteid amounts to about 15 grammes, and will be contained in 31 grammes of urea; 100 grammes of proteid contain about 50 grammes of carbon, while 31 grammes of urea contain only 6 grammes of carbon; so that in the translation of nitrogen from 100 grammes proteid to 31 grammes urea a margin of about 44 grammes carbon is left. This carbon may be disposed of by elimination in CO<sub>2</sub>, or temporarily stored as glycogen or as fat. We know from experimental data that glycogen and sugar can be formed from proteid, and that fat also can have its source in proteid (pp. 215, 258, 267).

(2) *The oxygen deficit*.—Comparing the composition of a carbohydrate (starch or sugar) with that of fat, we notice that the former contains oxygen and hydrogen in the proportion which gives water, viz. 2 H and 1 O, whereas on referring to the formulæ of fats we find that the hydrogen is greatly in excess, *e.g.* in palmitic acid we have 32 H and 2 O. This excess hydrogen in

fat must be oxidised in the body and escape as  $\text{H}_2\text{O}$ . It is accountable for the greater part of the deficit of oxygen in expired  $\text{CO}_2$  as compared with inspired  $\text{O}_2$ . 100 grammes of fat contain about 12 grammes of H and 11 grammes of O. The complete oxidation of 1 gramme H requires 8 grammes O; the complete oxidation of 12 grammes H requires 96 grammes O; so that the oxidation of 100 grammes fat must be supplemented by about 85 grammes O, or nearly 60 litres.

The same holds good of proteids, but in less degree. Referring to the percentage composition of proteids, we find that 100 grammes proteid contain about 7 grammes H and 21 grammes O; 7 grammes H will be oxidised by 56 grammes O, so that the complete oxidation of 100 grammes proteid must be supplemented by about 35 grammes O, or nearly 25 litres.<sup>1</sup> Thus the total amount of additional oxygen required for the oxidation of 100 grammes fat and of 100 grammes proteid will be 85 litres. This amount will go off as  $\text{H}_2\text{O}$ , and be accounted for by a deficit of  $\text{O}_2$  in the expired  $\text{CO}_2$ .

From these considerations it will be evident how the nature of food will affect the amount of oxygen that will be subtracted from external respiration and diverted to the oxidation of excess hydrogen in fat and in proteid, and thus influence the respiratory quotient  $\frac{\text{vol. of expired CO}_2}{\text{vol. of inspired O}_2}$ . Obviously, the less the subtraction of O by H, the greater the quotient; the greater the subtraction of O, the less the quotient. On a pure carbohydrate diet, the subtraction will be least and the quotient greatest. On a proteid, and *a fortiori* on a proteid *plus* fat diet, the subtraction will be greatest, and the quotient least—relations that are illustrated by the numbers already quoted on p. 133.

Comparing the result derived by calculation from food data with that obtained by direct observation of the respiratory gas exchange, we recognise that the two results correspond fairly well; from food we calculate that the oxygen supplement should be rather more than 80 litres—in the respiration numbers given on p. 132 we find an oxygen excess of 108 litres; this is quite as close a correspondence as could be expected, and if it is not closer still, it is because these numbers have been given rather high with a respiratory quotient rather low, *i.e.* .8; if with the same

<sup>1</sup> For simplicity's sake we have assumed *complete* oxidation in the body; if, however, we make allowance for incomplete oxidation, *e.g.* a discharge of 33 grammes urea, the oxygen supplement with 100 grammes proteid will be a little over 20 litres.



absorption of oxygen we had had a quotient =  $\cdot 85$ , the oxygen excess would have come out as 81 litres.

It may be asked whether the amount of water given off per diem is greater or smaller than that taken in ; to this the answer is that it is *theoretically* greater, although practically the excess is not to be detected—88 litres of  $O_2$  would go off in about 140 grammes  $H_2O$ .

**The balance of nutrition.**—*Import equals export.*—We have learned that the average daily discharge of a man weighing 70 kilos is—carbon 230 grammes, nitrogen 15 grammes, and it is obvious that while a man's weight and health remain constant this daily export must be made good by an exactly equal import of carbon and of nitrogen in food—*i.e.* an average daily diet must contain at least 230 grammes of carbon and 15 grammes of nitrogen. Such an allowance would be low, and the diet should be estimated on a somewhat more liberal scale, *i.e.* to contain 300 grammes of carbon and 20 of nitrogen. Still, so long as the weight and state of an animal remain constant, any increase of import must be balanced by an increased export, and under these circumstances 300 grammes of carbon and 20 grammes of nitrogen are necessarily excreted. If an animal gives off less carbon and less hydrogen than it takes in, it must be gaining weight ; if it gives off more carbon and more nitrogen than it takes in, it must be losing weight. The first event happens in a young growing animal, the second in a starving animal. Let us consider in further detail the possible modes of departure from the equipoised state of normal nutrition.

*Gain or loss of nitrogen signifies gain or loss of flesh.*—If during a given period the nitrogen import exceeds the nitrogen export, nitrogen must remain in the body. It does so in the form of proteid, *i.e.* as flesh. An animal under these conditions gains flesh. If the nitrogen import falls below the nitrogen export, nitrogen must be lost from the body ; the animal is losing flesh.

*Gain or loss of carbon signifies gain or loss of fat.*—If the carbon import exceeds the carbon export, carbon remains in the body. It does so in the form of carbohydrate or fat, probably as fat. If the carbon import falls below the carbon export, carbon is lost from the body ; the animal is losing fat.

In short, a gain or loss of *nitrogen* by the body signifies a gain or loss of *flesh* ; a gain or loss of *carbon* signifies a gain or loss of *fat*. And if the percentage composition of flesh and of

fat be known, it is easy to calculate how much flesh and how much fat correspond with each gramme of nitrogen and of carbon lost or gained. Flesh contains 3·3 per cent. of nitrogen ; 1 gramme of nitrogen signifies, therefore, 30 grammes of flesh. Fat contains 75 per cent. carbon ; 1 gramme of carbon signifies, therefore,  $1\frac{1}{3}$  gramme of fat.

To what extent do observed facts bear out these theoretical conclusions ? The answer to this question is derived chiefly from experiments on the dog : the reason for this choice being that the animal is carnivorous, and can be kept in health during a

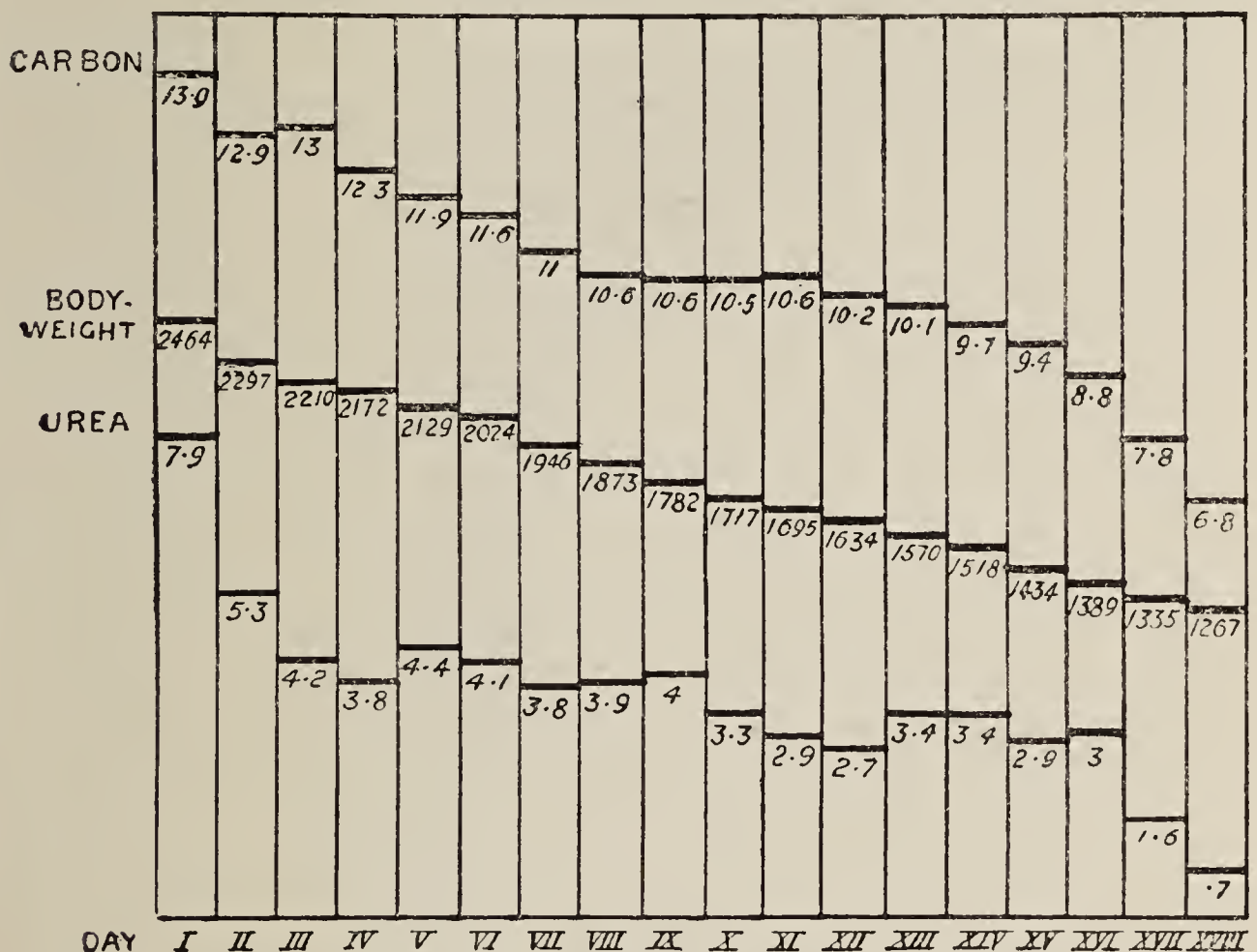


FIG. 108.

Graphic representation of the declining weight and of the daily excretion of a starving cat. (Data by Bidder and Schmidt.)

considerable period upon an exclusively proteid diet (meat); other advantages are, that a dog can be trained to perform definite amounts of work on a treadmill, and to give up the urine either naturally or by catheter when required to do so, and that the error introduced by neglecting losses of nitrogen otherwise than by the kidney does not amount to more than 1 to 2 per cent. of the total nitrogen. On man, the amount of nitrogen eliminated in the faeces amounts to about 10 per cent. of the total discharge ; on herbivora, half the total nitrogen, or even more, may be contained in the faeces.



**Starvation.**—*Export is greater than import.*—An animal (*e.g.* a dog or cat) receiving no food, receives no nitrogen and no carbon, yet continues to excrete urea and carbonic acid until the end of its life. In this case there is obviously no equilibrium of nitrogen or of carbon, but an unbalanced expenditure of both. The animal loses flesh, loses fat, and loses weight. The incidence of these losses is well illustrated in the accompanying diagrams. In fig. 108 are given the amounts of carbonic acid and of urea excreted, and the body-weight from day to day of the starvation period. In fig. 109 are given the total loss per cent. of the most important organs and tissues at the end of starvation.

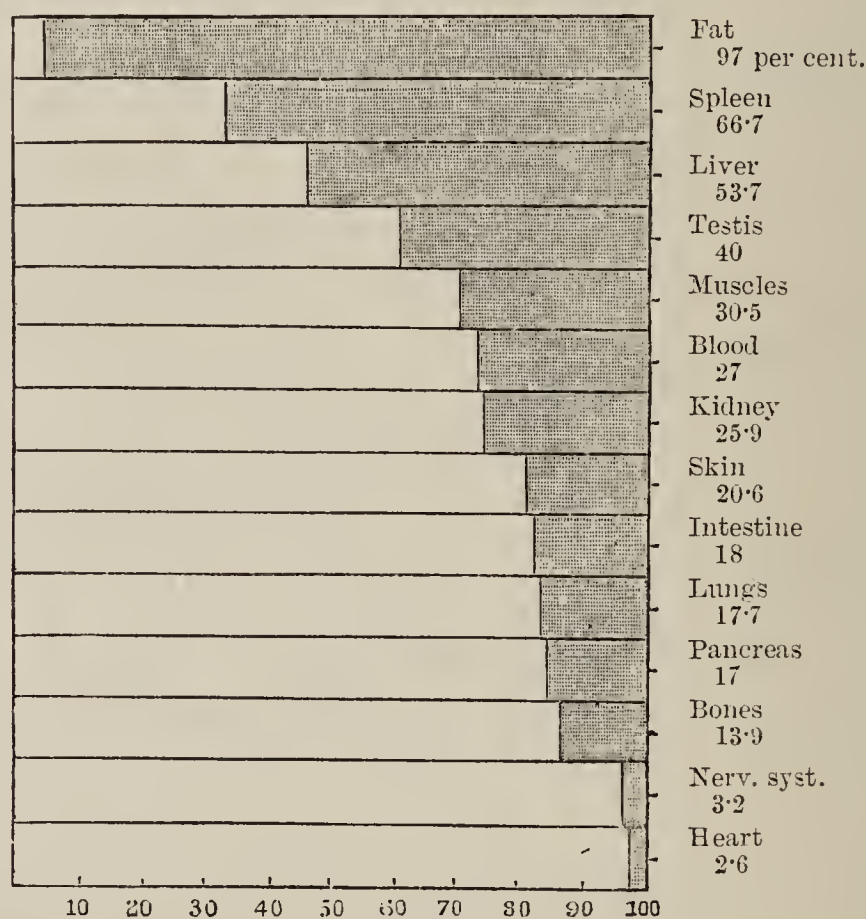


FIG. 109.

Graphic representation of the percentage of different tissues lost during starvation; the shaded areae represent loss, the unshaded areae residue at death. (According to Voit's analyses.)

*Minimum nitrogen requisite to nitrogenous equilibrium with an exclusively proteid diet.*—It might be expected that, if nitrogen in proteid food equal in quantity to nitrogen excreted during starvation were administered, the loss of nitrogen would be balanced. The expectation would not be confirmed by the event, for the animal would secrete all the more nitrogen, and would require about three times this amount, *i.e.* three times the amount of the starvation export. Then only would import balance export and the animal be in nitrogenous equilibrium. Beyond this minimum

amount, a further increase of food-nitrogen would still be balanced by a corresponding increase of excreted nitrogen, and the animal would continue to be in nitrogenous equilibrium. Excessive amounts of meat would be liable to cause diarrhoea, and thus put an end to the observation. But from 'minimum' to 'excessive' the range is considerable; nitrogen equilibrium is maintained with nitrogen imports of considerable magnitude; this is the normal state of a healthy or even of a voracious man, his nitrogen import is far above the minimum necessary, but is balanced by a correspondingly large nitrogen export, and he remains in nitrogenous equilibrium, with perhaps a breakdown of his kidneys or an attack of gout close at hand. Exact experimental results relating to nitrogenous equilibrium are, however, possible only on carnivorous animals—usually dogs.

*Example.*—A dog weighing 10 kilogrammes excretes during the first days of starvation about 5 grammes of nitrogen per diem. This amount of nitrogen would be contained in about 150 grammes of meat. In accordance with what has been said above, a daily allowance of 150 grammes meat, containing only just as much nitrogen as is contained in the starvation export, would be insufficient to balance the daily loss. The excretion of nitrogen would be increased, and would be in excess of the nitrogen income of 5 grammes derived from 150 grammes of meat. It would be between 5 and 10 grammes, and the animal on the daily allowance of 150 grammes of meat would continue to lose flesh and eventually starve. Nitrogenous equilibrium would not be reached until nearly 500 grammes of meat are given per diem, and now the import and export of nitrogen would each be about 15 grammes. As already stated above, nitrogenous equilibrium is not obtained until the food contains about three times the amount of nitrogen excreted when no food is given, and in the case of a carnivorous animal fed upon meat, the daily ration must amount to about  $\frac{1}{20}$  the body-weight of the animal. Beyond this minimum (500 grammes meat) the animal is capable of disposing of an additional quantity, still remaining in nitrogenous equilibrium: 1,000 grammes may be the daily ration consumed; in this case the nitrogen of the food and the nitrogen of the excreted urea will each amount to about 30 grammes per diem.

*Nitrogenous deficit.*—In the above example the nitrogen income and outgo balance each other; there is no deficit on either side. But in many experiments the two quantities have been



found not to balance ; the amount of nitrogen exported in urea has been found to fall short of the nitrogen imported in proteid. The deficit has been accounted for as being due to the discharge of nitrogen in a gaseous state by respiration. The fact has, however, not been proved—the amount of  $\text{NH}_3$  in expired air is infinitesimal, about .001 grm. per hour—and we have against it the observations of nitrogenous equilibrium with varying quantities of proteid food. It appears most probable that the so-called nitrogenous deficit is only apparent, and due to a deficit of analysis.

Returning to our example, let us consider the carbon equation and the body-weight of the dog weighing 10 kilogrammes. During the total deprivation of food the dog is obviously losing flesh, losing fat, and losing weight. With the insufficient allowance of 150 grammes of meat the dog is still suffering the same losses, but less rapidly. With the sufficient diet of 500 grammes, the dog is neither losing nor gaining flesh, but may be losing or gaining weight. 500 grammes of meat contain nearly 70 grammes of carbon, and it is possible that the respiration of the animal is such that more or less than this amount is exported as carbon dioxide—if more, the animal is losing fat and losing weight ; if less, the reverse is the case. With the more than sufficient allowance of 1,000 grammes, and the animal in nitrogenous equilibrium, it is probable that the carbon export will not amount to the carbon import, and the animal will consequently be putting on fat and gaining weight. The value of this increase, and in the case of the 500 grammes allowance, whether there is increase or decrease, is to be ascertained by measuring the amount of  $\text{CO}_2$  excreted per diem.

These statements will be made clear by considering the accompanying balance-sheet, which gives the nitrogen and carbon with inadequate, adequate, and more than adequate diets of meat given to a small dog. The excretion of nitrogen otherwise than in urea is omitted from consideration as being a negligible quantity.

*Influence of age.*—The case of a dog on a diet of 500 grammes of meat is deserving of some further consideration. The amount is that theoretically the smallest adequate to the maintenance of a 10-kilogramme dog in nitrogenous equilibrium. It would be found that other things being equal, a young dog would require a higher minimum than an old dog, and we recognise in this fact an experimental illustration of what experience shows to be

DOG WEIGHING 10 KILOGRAMMES<sup>1</sup>

Daily income	N	C	Daily expenditure	N	C	Remarks
I. None	0	0	Urea 10·7 grammes CO <sub>2</sub> 110 grammes Total . . .	5 0 5	2 30 32	The dog is starving rapidly, losing per diem 5 grammes N and 32 grammes C, equivalent to 150 grammes flesh and 43 grammes fat.
II. Meat 150 grammes	5	20	Urea 15 grammes CO <sub>2</sub> 143 grammes Total . . .	7 0 7	3 39 42	The dog is still starving, losing per diem 2 grammes N and 22 grammes C, equivalent to 60 grammes flesh and 30 grammes fat.
III. Meat 500 grammes	16·6	66·6	Urea 35·6 grammes CO <sub>2</sub> 220 grammes Total . . .	16·6 0 16·6	7 60 67	The dog is in nitrogenous equilibrium, neither gaining nor losing flesh. The dog is also with the above carbon values neither gaining nor losing fat. It would, however, be found in an actual experiment that, whereas the nitrogen export would remain practically constant and in equilibrium with import during many days, the carbon export would vary considerably, implying, if above 67 grammes, a loss of fat, if below 67 grammes, a gain of fat.
IV. Meat 1,000 grammes	33	133	Urea 70 grammes CO <sub>2</sub> 275 grammes Total . . .	33 0 33	14 75 89	The dog is still in nitrogenous equilibrium, neither gaining nor losing flesh, but gaining fat at the rate of 58 grammes per day (44 grammes C). The animal is gaining weight with this deposition of fat. The experiment is likely to terminate with an attack of diarrhoea, an event which would be retarded by allowing the dog plenty of exercise.

<sup>1</sup> According to Senator's measurements, the average daily heat-production by a dog of this weight in the fasting state would be about 500 kilocalories. The heat-value of meat=about 1 kilocalorie per gramme, of fat=about 9 kilocalories per gramme.



requisite in the feeding of children, who require a larger proportion of proteids than do adults, taking into account the difference of body-weight. Thus, the mixed diet of an adult should contain 2 grammes of dry proteid per kilogramme body-weight, while that of an infant one year old should contain double that amount, viz. 4 grammes per kilogramme body-weight. The *state of nutrition* is also an influential factor; other things being equal, an ill-nourished animal is accommodated to and requires per diem a smaller allowance than a well-nourished animal, and the same animal passing from a well- to an ill-nourished state gradually accommodates itself to a lower minimum, while in passing from an ill- to a well-nourished state it requires an increasing minimum allowance, and becomes able to tolerate and dispose of a larger excess above that minimum. We recognise in these facts experimental illustration under simple conditions of what takes place in man under more complex conditions; many poor persons are accommodated to a much lower minimum than that which would be necessary to keep a well-to-do person in good condition; many well-fed persons are accommodated to a much higher minimum, and can tolerate an excess diet far greater than could be suddenly disposed of by an ill-fed individual. Such persons, however, run the risk of becoming overloaded with uric acid, or of overworking their kidneys; they are liable to be purged of undigested food by diarrhoea, or they find it necessary to habitually stimulate a 'sluggish' digestion, and exercise, instead of being the effect of natural inclination, becomes to them a hygienic observance.

*Peptone=proteid.*—The experiment has been made of substituting peptones for undigested proteid in the feeding of dogs. It has been found that peptones are capable of taking the place of proteid, and that dogs fed with peptones alone can grow and fatten and be kept in nitrogenous equilibrium.

*Effects of fat, carbohydrate and gelatine, added to proteid.*—If in addition to proteid an experimental diet is made to include fat or starch or gelatine, it will be found possible to establish nitrogenous equilibrium with a smaller quantity of proteid than when the latter is administered alone. In the case of the 10-kilogramme dog requiring per diem at least 500 grammes of meat to be brought into nitrogenous equilibrium, the result would be obtained with half that quantity of meat *plus* 50 grammes of fat. Each of the above-named substances—fat, carbohydrate, and gelatine—is a natural ally and companion of proteids in normal diets,

although administered alone none of these substances can support or even prolong life; an animal fed with fat, or with carbohydrate, or with gelatine, or with all these substances *sine* proteid, will speedily die of starvation.

*Effect of proteid upon fat.*—Fat alone cannot support life; added to proteid, it economises the latter, as above stated. As regards the converse relation, proteid administered with fat accelerates the consumption of fat; similarly proteid administered in considerable quantity to a fat animal accelerates the consumption of fat, and the animal loses fat. The Banting system of treating obesity by a diet containing abundant proteid and restricted fat and carbohydrate, is based upon this principle, which is also acted upon in athletic training; and the recent observations of Pflüger are to the same effect, viz. that tissue-change is intensified by proteid, retarded by fat. We must, nevertheless, remember that fat in the body may have proteid in the food as an antecedent, although the chief relation between the two is that proteid accelerates expenditure of fat while fat economises or obstructs the expenditure of proteid.

*Effect of carbohydrate upon fat.*—The popular notion that ‘sugar is fattening’ appears to be borne out by experiment. There can be no doubt that carbohydrates do promote the fattening of cattle; the analyses of Lawes and Gilbert have placed this point beyond doubt. These observers estimated the amount of fat put on by cattle during a given time, in comparison with the amount of fat contained in the food consumed during the same time, and found that the amount of fat gained was more than four times as great as the amount of fat contained in the food. Whence could the fat be derived? It might have been from proteid, or it might have been from carbohydrate, and it cannot be positively stated from which of the two sources it proceeded. This much is certain, that carbohydrate was essential to the fattening, but it is an open question whether it produced this result (1) by direct conversion of carbohydrate into fat, or (2) by economising the combustion of body-fat laid down from accompanying proteid. The first alternative is generally regarded as the more probable, and, although as yet it is not an undisputed conclusion, the weight of evidence is decidedly in its favour. It is to be remembered that, although carbohydrates fatten, they cannot entirely replace fat in the diet of man.

*Effect of fat upon carbohydrates.*—Fat in the food, if it enters



the body, is oxidised at once, or holds its place as fat. There are no observations to show that it can become or give rise to either proteid or carbohydrate in the body. All observers are agreed that fat does not assist in forming glycogen in the liver; Seegen, indeed, states that fat may directly form sugar in the liver, but his observations are unconfirmed.

**The normal diet of man.**—It is best, and, indeed, absolutely necessary, that human food should include *proteids*, *fats*, and *carbohydrates*, to which must be added *salts*, *oxygen*, and *water*. The absence of any one of these proximate principles kills by starvation. Absence of oxygen kills in two or three minutes, absence of water in a few days, absence of proteid, or fat, or carbohydrates in two to three weeks, and the absence of salts cannot be endured for a much longer period. That common salt is one of the necessities of life is attested by the risks which men have run to obtain it when scarce, and by the ‘salt-licks’ frequented by wandering troops of animals.

How little can a healthy man endure life upon, and how much must be given to him if he is to be kept efficient? The answer is afforded by the dietaries of armies, prisons, and gangs of navvies. In round numbers, and omitting circumstantial considerations, the minimum daily income of a man in full work must be about 5 per cent. of his body-weight, viz.

Solid food	.	.	.	.	.	1 per cent.
Oxygen	.	.	.	.	.	1 „
Water	.	.	.	.	.	3 „

That is to say, a man of 70 kilogrammes must get

Solid food	=	700 grammes
Oxygen	=	700 „
Water	=	2,100 „
<hr/>		
Total	=	3,500 „

The 700 grammes of solid food should contain about

140 grms. proteid
105 grms. fat
420 grms. carbohydrate
35 grms. salt.

Or, to reduce those numbers to an easily remembered form, from which the adequate diet of men of different weights can be estimated, it should contain per one kilogramme body-weight,

2 grms. proteid ; 1·5 gm. fat ; 6 grms. carbohydrate ; 0·5 gm. salt—in all 10 grammes, or *one per cent.* of solid food.

These numbers are applicable to normal adults doing ordinary work. The estimate is a liberal one, especially as regards the proteid. Voit gives as adequate :—Proteid, 118 to 137 ; fat, 56 to 117 ; carbohydrate, 352 to 500. Ranke's 'normal diet' is :—Proteid, 100 ; fat, 100 ; carbohydrate, 240 grms.

The chief conditions that will modify the amounts of an adequate diet are *work, health, and age.*

*Effect of work.*—A person doing no work at all will subsist upon amounts somewhat smaller than those given above ; a person engaged in exceptionally heavy work (soldier in the field, prize-fighter, navvy, &c.) will require larger amounts :

No work . . . . .	6·6 grs. solid food per 1 kg.
Ordinary work 100·000 kgm. . . . .	10 „ „
Hard work 200·000 kgm. . . . .	15 „ „

*Effect of food upon excretion of urea.*—The amount of nitrogen excreted in urea is directly dependent upon the amount of nitrogen taken in proteid. A dog taking 500 or 1,000 grammes of meat eliminates 35 or 70 grammes of urea, and if one day he takes and digests double the amount of proteid, he will during the next day excrete about double the amount of urea. The condition of the dog or man is one of nitrogenous equilibrium, or, in other words, the amount of urea excreted is in direct relation with the amount of eaten and digested proteid.

*Effect of muscular exercise upon excretion of urea.*—It might be expected that great muscular exertion should be attended with a disintegration of muscle and a corresponding increase in the amount of urea excreted. The expectation has not been verified by experiment. Muscular exertion *per se* gives rise to no marked increase of urea excreted ; in other words, there is no demonstrable relation between amount of work done and amount of urea excreted. The best-known observations in justification of this statement are those of Fick and Wislicenus upon themselves, and those of Flint and of Pavy upon a pedestrian. But while we must admit that the discarded doctrine of a ratio between work and wear of muscle, as evidenced by urea, is incorrect, we must also admit that the contradictory doctrine has been expressed too absolutely. Recent experiments show that urea excretion does vary a little with muscular action, although the variations are so small as to be easily swamped and hidden by the larger fluctuations due to variations of diet ; Zuntz's observations are to the



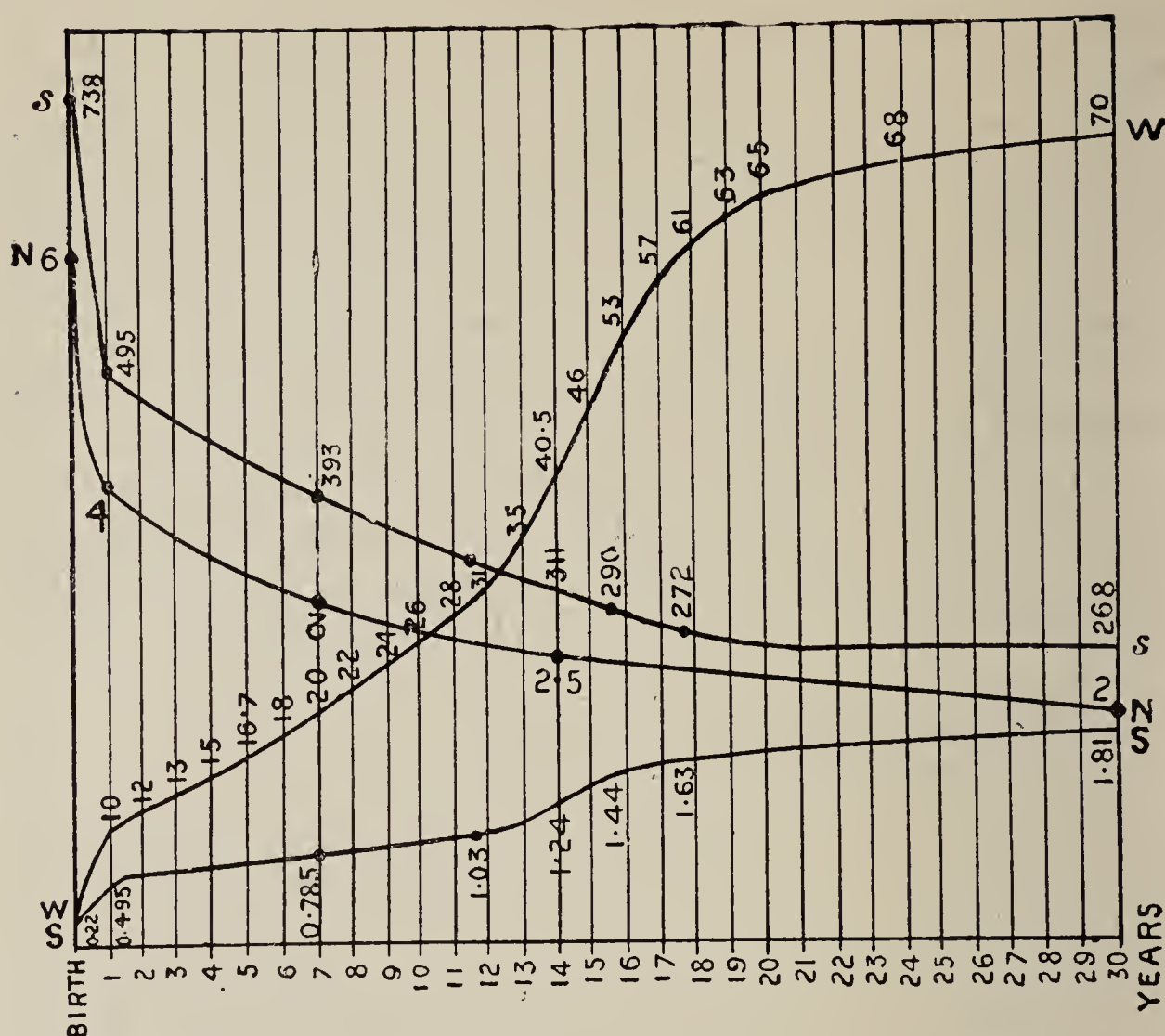


FIG. 110.—WEIGHT AND SURFACE OF BODY AT DIFFERENT AGES. PROTEID REQUIRED.

W W = body-weight in kilos (1 mm. = 1 kilo).

S S = body-surface in sq. meters (1 cm. = 1 sq. meter).

s s = body-surface per 1 kilo body-weight in sq. cms. (1 mm. = 10 sq. cms.).

N N = proteid per kilo body-weight in grammes (1 cm. = 1 gm.).

The surface measurements are calculated upon the assumption that the surface of an animal bears a constant relation to its weight, such that

$$\frac{S}{(\text{in square centimeters})} = 10.5 \sqrt[3]{W^2} \quad (\text{in grammes}).$$

The ratio generally adopted is rather higher, viz. :  $S = 12 \sqrt[3]{W^2}$ .

effect that muscular exercise, if excessive—*i.e.* if pushed to the point of producing dyspnœa—causes disintegration of organ-proteid, and increased discharge of urea; and Dubois' observations on the loss of weight suffered by marmots during hibernation (20 per cent. in 160 days of sleep versus 20 per cent. in 12 days of active wakefulness) clearly show that consumption of tissue is entailed by normal or even subnormal activity. That such consumption is inclusive of proteid tissue is proved by the urea excretion which proceeds during hibernation in the absence of proteid food.

Although muscular exercise may produce little or no discoverable increase of the nitrogen discharge if the nitrogen income is maintained constant, it may, under ordinary conditions of life,

give rise to a very considerably increased discharge, by provoking a greater ingestion of proteid. In an animal fed upon proteid alone the total energy evolved as heat and as work must ultimately have its source from proteid, and whether or not that proteid forms carbohydrate in its passage through the body is a matter of secondary importance. Pflüger's dog, *e.g.* weighing 30 kilos, was in equilibrium on a pure flesh diet of 2 kilos per diem when doing no work, and required a daily supplement of .5 kilo when doing work estimated at 109·608 kgm. per diem; in this case the work must have been sustained from the proteid, and, comparing the heat-values of the two quantities, the yield of work comes out as equivalent to between 40 and 50 per cent. of the supply of potential energy afforded by the supplementary food.

*Diet during infancy.*—A child or infant requires absolutely less food than a full-grown man, but relatively to their respective body-weights a much larger quantity. Thus, it would not be sufficient to give to a child of 7 or 14 kilos only  $\frac{1}{10}$  or  $\frac{1}{5}$  the amount required by a man of 70 kilos, and we should find in drawing up a diet-table for children at different ages, that the area of the body-surface is a far better proportional indicator than the body-weight. This is especially the case as regards proteid, and will be evident on examination of the table below.

Age	Body-weight	Body-surface	Total dry proteid per diem	Do. per kilogramme body-weight	Do. per sq. meter body-surface
	kilogrammes	square meter	grammes	grammes	grammes
Birth	3	0·22	18	6	90
1 year	10	0·5	40	4	88
7 years	20	0·8	62	3	87
14 „	40	1·24	100	2·5	88
30 „	70	1·8	140	2	85

The last two columns of this table show that, in proportion to body-weight, the amount of proteid required by an infant is greater than that required by an adult, while in proportion to body-surface it is approximately constant (*viz.* between 80 and 90 grammes per square meter). As regards other constituents, the best guide to their amounts is the composition of milk, which is normally the only food required by infants during the first months of life. In round numbers human milk contains per 100 c.c.

3 grammes proteid  
 3·5 „ fat  
 5 „ sugar



An infant at the breast must therefore take per diem 500 to 1,000 c.c. of milk to get

15 to 30	grammes	proteid,
20 to 40	„	fat,
25 to 50	„	carbohydrate.

If artificial feeding is required, cow's milk is usually first resorted to, and seeing that cow's milk is, as compared with human milk, richer in fat and proteid, poorer in sugar, it is necessary to dilute it and to add sugar to it. A common error in the feeding of infants is the premature augmentation of the carbohydrate element, 60 to 90 grammes carbohydrate being the amount commonly given in diet tables to correspond with 20 to 36 grammes proteid. The composition of milk is, however, such as to indicate that the carbohydrate should not largely exceed the proteid constituent, during, at any rate, the first 9 months of life. After this period the requirements may be regarded as approximating towards those of the adult, in whom, as we have seen, the amount of carbohydrate required is three times that of proteid.

A growing infant excretes less carbon and less nitrogen than it takes in; it gains in weight and puts on both flesh and fat; during the first year of its life it gains about 7 kilos. Taking half of this amount to be flesh, this would mean a gain of about 120 grammes nitrogen, *i.e.*  $\frac{1}{3}$  gramme per diem; the average proteid income, 30 grammes, contains about  $4\frac{1}{2}$  grammes nitrogen; the amount of nitrogen kept back in the flesh of a growing infant up to the end of the first year of life is thus about  $\frac{1}{13}$  of the nitrogen contained in the total proteid consumed; during the second year it is about  $\frac{1}{60}$ , during the third about  $\frac{1}{140}$ .

*Diet during lactation.*—Consider the case of the mother, supposing her to weigh say 50 kilos, and to receive for her own needs alone an average proteid allowance of 100 grammes. If she is to supply a child with one litre of milk containing 30 grammes of proteid per diem, she should receive an addition of about 90 grammes to her food, *i.e.* her day's proteid should amount to nearly 200 grammes, a requirement that is recognised in the instructions issued in France to persons employing a wet-nurse, to the effect that she must receive nearly twice as much food as an ordinary woman. While nursing she will also excrete considerably more urea than in her normal state.

That so large an increase in the proteid allowance should be

requisite during lactation is no less apparent if we consider the facts already alluded to concerning the relation between proteid and fat. We have seen that proteid can fatten, that 100 grammes proteid contain about 44 grammes of surplus carbon, which may go to form fat or carbohydrate; now it is shown both by experience and by experiment that not fat but proteid is the substance from which milk-fat is formed. The day's milk yielded by a wet-nurse contains, say, 30 grammes of proteid and 40 grammes of fat; the proteid supplement to her ordinary diet is, say, 90 grammes; of this amount 30 grammes are accounted for in the milk; the remaining 60 grammes contain 9 grammes nitrogen (which will go off in about 20 grammes urea) and 27 grammes surplus carbon (which might form part of 36 grammes fat). This is as good an illustration as can be desired, showing that the proteid molecule can split into a nitrogenous moiety, urea and a non-nitrogenous moiety, fat. In the case just considered 90 grammes of food proteid are accounted for by 30 grammes of milk proteid + 37 grammes of fat + 20 grammes of urea.

*Alterations of nutrition in disease; diabetes; fever; the convalescent state.*—The food requirements of the body are, as a rule, much diminished in disease; the activity of the tissues is depressed, little or no external work can be done, and diet must be calculated on a much lower scale.

There are, however, certain exceptions to the general rule; in *diabetes*, the amounts of food and of water that are 'consumed' are sometimes enormous; but is food really consumed in a physiological sense, that is to say, utilised in the nutrition of tissue? The wasting and loss of body-weight, the excessive discharge of sugar and of water, and, in certain cases, the degeneration of proteid into fat, and, finally, the incapacity of diabetic subjects to perform work, all go to show that the body is then incontinent of food, which its tissues fail to assimilate.

In *fever*, as a rule, we find that much less solid food is consumed, while the demand for water is increased. Here, again, the tissue-waste is increased, morbid products are formed, and the increased demand for water may be viewed as the attempt of the organism to wash itself clean; copious perspiration is a sign of success, and a token that the critical point has been turned. In *convalescence* from fever we have a true instance of increased nutrition; the tissues have wasted during invasion, and when released from the oppressive agencies reassert their want of food to an enhanced degree; 'appetite' is good, more food is taken



by the mouth and consumed in the tissues, the body-weight rises again and strength returns; the chief danger is that the renewed appetite of the tissues should overtax the digestive organs, which have shared in the general depression.

**Hypertrophy and atrophy of parts and of organs.**—If a limb or organ is more than usually exercised, it grows larger than usual; if it is less than usually exercised, it dwindles and becomes smaller than usual. The first event is called hypertrophy, *i.e.* over-nutrition; the second, atrophy, *i.e.* deficient nutrition; and these effects, which we can see, and touch, and trace back to evident causes, give us an insight into the normal process that we could not otherwise obtain. A part that is more used gets more 'food'; its vessels dilate, and the blood-flow is increased in response to the necessities of the tissues. A part that is less used, gets less food; its vessels contract, and the blood-flow is diminished. In either case the supply is adjusted to the demand through the agency of afferent and efferent nerves. Experimental and pathological illustrations of both processes are abundant. If the cervical sympathetic of a young rabbit be divided, the vessels of the ear on that side dilate, more blood passes through the part, and the ear grows faster and larger than on the other side. The skin round an old-standing sore may become thicker and more hairy in consequence of the increased blood-supply in the neighbourhood of the irritation. The spur of a cock transplanted from the leg to the comb grows to an extraordinary degree, because it gets more blood than usual.

Instances might be multiplied. Any hollow viscus grows larger and thicker if the channel from it is narrowed; any limb or single muscle grows bigger if its daily work is increased, on the principle—more work, more food, more tissue. Hypertrophy of the heart, of the bladder, of the intestines, the disproportionately well-developed legs of a ballet-dancer, or arms of a blacksmith, the contrast between the arms of a navy and those of a clerk, all illustrate this principle. The converse effects are no less marked, though, perhaps, less striking and apparent; the consequences of atrophy or deficient nutrition are to be seen in every old person, in less active organs, less work, less active circulation, in dwindling and degenerating tissues.

That food-supply by blood to tissue is controlled and measured out by nerves, we have already learned from the study of vaso-motor action; whether the intimate processes of tissue-nutrition

are governed by nerves, we shall inquire when we come to study the obscure question of *trophic nerves*.

**Construction and calculation of dietaries.**—The proximate principles—proteids, fats, and carbohydrates—convey into the body the necessary nitrogen and carbon, and are themselves conveyed into the body in the various food-stuffs or articles of diet in common use. Without entering into any minute description of the properties and qualities of all the articles of food that may be named on a bill-of-fare, we will very briefly consider one or two staple food-stuffs, and point out how diet-tables should be made use of in order to calculate an adequate food allowance, *i.e.* an allowance not far short of **20 grammes nitrogen** and **300 grammes carbon**.

*Food-stuffs.*—It should be said in the first place that the proteid or nitrogenous principles can be derived from animal and from vegetable sources; that, of the non-nitrogenous principles, fats are also derived from both sources, while carbohydrates are of entirely vegetable origin, with the exception of milk-sugar and of glycogen.

APPROXIMATE COMPOSITION OF SOME COMMON ARTICLES OF DIET.

Food-stuffs	Proximate principles				Elements	
	Water per 100	Proteid per 100	Fat per 100	Carbo- hydrate per 100	Carbon per 100	Nitrogen per 100
Milk . . .	86	4	4	4	7	0·6
Butter . . .	7	1	92	—	70	0·15
Eggs . . .	75	14	10	—	15	2
Beefsteak . . .	70	22	5	—	15	3·3
Bread . . .	40	8	1·5	50	27	1·25
Potatoes . . .	75	2	—	21	10	0·3
Oatmeal . . .	15	12	5	65	40	2
Dried peas . . .	15	22	2	60	40	3·3
Rice . . .	10	5	1	83	40	0·75
Cocoa powder . . .	10	15	50	25	55	2·2
Cheese . . .	40	35	25	—	35	5·2
Beer . . .	90	1	—	10	5	0·15

Referring to the table, the following points are to be remarked. The chief nitrogenous food-stuff is meat, but many vegetables, *e.g.* peas, are not inferior to meat as regards the percentage of nitrogen they contain. This does not imply, however, that the nutritive value of peas is equal to that of meat; most people will absorb 3·3 grammes of nitrogen more easily from 100 grammes of meat than from 100 grammes of split peas. All the nitrogenous food-stuffs contain carbon as well as nitrogen; this we



already know from the composition of proteids, which contain about 50 parts of carbon to each 15 parts of nitrogen. Of the 'non-nitrogenous' food-stuffs, only butter, which is the fat of milk, and the ordinary fat that is attached to meat, are literally non-nitrogenous. Vegetables, such as potatoes, are the chief source of the non-nitrogenous carbohydrate, but they are not non-nitrogenous food-stuffs, inasmuch as they also contain nitrogen. Bread in particular, the most important of all food-stuffs, contains nitrogen as well as carbon.

The desirable proportion between nitrogen and carbon in a diet is accepted to be 1 N to 15 C by weight. In meat the proportion is about 1 N to 5 C; so that if a person should attempt to live upon meat only, he must swallow too much nitrogen if he is to get a right amount of carbon, or too little carbon if he is to get a right amount of nitrogen. In bread the proportion is about 1 N to 22 C; so that if a person should be restricted to bread only, he must swallow too much carbon if he is to get a right amount of nitrogen. But while neither of these food-stuffs alone yields the right proportion, both together will do so, and in ordinary life meat and bread are taken together, these two food-stuffs constituting the foundation of every prosperity diet, to which they contribute nearly all the nitrogen and three-fourths of the carbon. Theoretically, indeed, it would be possible to obtain exactly the right amounts of carbon and of nitrogen by duly adjusted amounts of these two staple articles, meat and bread, but practically it is necessary to take into account a due proportion of fat, which has by experience been proved indispensable to the maintenance of health, and not fully replaced by carbohydrate. Considerations of cost and of taste will also have to be taken into account.

*A day's diet.*—As the foundation of a day's diet we may put down

	Carbon	Nitrogen
Bread, 1 pound (450 grammes) . .	117 grms.	5.5 grms.
Meat, $\frac{1}{2}$ pound (225 grammes) . .	34 „	7.5 „
Making a total of . .	151 „	13 „

The chief deficiency here is of the carbon; nitrogen is also defective, but less so, though not less so than in many a necessity diet; we are, however, at this moment constructing the diet of a properly fed, healthy, active man.

Theoretically as well as practically the foundation from which we have started will be completed by fat (with the meat, or as

bacon or as butter); from all sources about  $\frac{1}{4}$  lb. (112 grammes) containing 84 grammes carbon.

The diet is now

	Carbon	Nitrogen
Bread, 1 pound . . . . .	117 grms.	5·5 grms.
Meat, $\frac{1}{2}$ pound . . . . .	34 „	7·5 „
Fat (butter, bacon, &c.), $\frac{3}{4}$ pound . . . . .	84 „	—
Making a total of . . . . .	235 „	13 „

This foundation is ordinarily supplemented or in part replaced by a variety of articles, the most important of which are *potatoes*, *rice*, *eggs*, *milk*, *cheese*, *oatmeal*, *peas*, *beans*, and *cocoa*.

*Potatoes* to a certain extent replace bread and butter; they are especially a carbon supply, also—but in a very low proportion—a nitrogen supply. One pound of potatoes (450 grammes) contains 45 grammes carbon and 1·3 nitrogen. An Irish labourer consuming 10 pounds of potatoes per diem, and nothing else, would get 450 grammes carbon and 17 grammes nitrogen, but he would be ‘ill-fed.’

*Cheese* is a very prevalent substitute for meat among agricultural labourers;  $\frac{1}{8}$  lb. cheese (56 grammes) contains 20 grammes carbon and 3 grammes nitrogen. *Half a pint of milk* (283 c.c.) contains about 20 grammes carbon and nearly 2 grammes nitrogen. *Two eggs* (weighing about 100 grammes) contain about 15 grammes carbon and 2 grammes nitrogen.

*Peas*, *beans*, and *lentils* are to be mentioned as forming the backbone of a vegetarian diet, in which they take the place that meat occupies in ordinary diet, forming the chief nitrogen supply. *Rice* is here mentioned as being a staple article of food among Eastern nations; *oatmeal* on account of the important place it holds in the dietary of North Britain. But, seeing that all these articles are accidental rather than habitual items in ordinary English life, we shall not take them into further account in this connection.

Adding some of the preceding accessories to our fundamental diet we have as an ideal complete diet:—

	Carbon	Nitrogen
Foundation	1 pound bread . . . . .	117 grms.
	$\frac{1}{2}$ pound meat . . . . .	34 „
	$\frac{1}{4}$ pound fat . . . . .	84 „
Accessories	1 pound potatoes . . . . .	45 „
	$\frac{1}{2}$ pint milk . . . . .	20 „
	$\frac{1}{4}$ pound eggs . . . . .	15 „
	$\frac{1}{8}$ pound cheese . . . . .	20 „
	335 „	21 „



This is a liberal diet with a margin above the normal, but it is somewhat costly, viz. at ordinary market prices nearly one shilling.

More detailed consideration of diet belongs to practical medicine; all that is desirable from a physiological standpoint is to realise certain fundamental principles and certain prominent illustrations of these. It is important to become familiar with the use of tables of food analysis, with clear knowledge of the essential points that should be looked for. The short table used above in the construction of a normal diet gives in round numbers the compositions of the commonest food-stuffs rated in proximate principles and in their two important elements of income. We may use this table again to estimate the nitrogen and carbon values of a hospital 'ordinary' diet, *e.g.* :—

	Carbon	Nitrogen
2 pints of tea (with sugar) (1,134 c.c.) . . . . .	? 25 grms.	— grms.
$\frac{1}{2}$ pint of milk (283 c.c.) . . . . .	20 „	1.7 „
1 pint of beef-tea (567 c.c.) . . . . .	? 7 „	? .6 „
4 oz. cooked meat (113 grms.) . . . . .	20 „	4 „
$\frac{1}{2}$ lb. potatoes or vegetables (225 grms.) . . . . .	22 „	.7 „
$\frac{3}{4}$ lb. bread (339 grms.) . . . . .	75 „	4 „
$\frac{3}{4}$ oz. butter (21 grms.) . . . . .	16 „	— „
Sum-total . . . . .	185 „	11 „

Obviously, this sum-total must have been inadequate to sustain life in a large percentage of cases, especially if the patients were not confined to bed and entering upon convalescence, and as a matter of course it is supplemented by the liberal allowance of 'extras.' We may add for the sake of comparison the following figures :—

'Starvation diet' (East-end needle-woman) . . . . .	Carbon	Nitrogen
	200 grms.	9 grms.
'Almost starvation diet' . . . . .	264 to 270 „	11 to 13 „
'Light-labour' prison diet . . . . .	295 „	14 „
'Hard-labour' prison diet . . . . .	335 „	16 „
Fairly nourished operatives . . . . .	390 „	22 „
Military prison diet . . . . .	340 to 400 „	18 to 20 „

All these diets are too low to sustain life at a normal level of energy for any prolonged period in the various classes of persons to whom they are administered. Flagging energy and loss of flesh are quickly produced in the first three examples, and even in the last three it has been found necessary to raise the diet or to lower the labour.<sup>1</sup>

<sup>1</sup> The veil that measurements in pounds, ounces, pints, and grains casts over a diet-table is best removed by converting them at once into grammes and cubic

*The factors of nourishing value.*—To properly estimate the nourishing value of a food we must take into account, not merely its percentage of carbon and nitrogen, but also its palatability and its digestibility. A 'nourishing' food must not only be chemically adequate, it must also be palatable and digestible. It may possess any one, or even two, of these three essentials, and yet not be nourishing. Thus, dried peas and beans, or even horse-chestnuts and acorns, may be chemically adequate, but they are not palatable. Cheese and hard-boiled egg stand high as regards chemical value, but are very indigestible. Conversely, it may happen that a substance is particularly digestible, so that it has a nourishing value which its chemical composition would not have led us to expect; gelatine, for instance, in the form of calves'-foot jelly, is of indisputable practical value in the nourishment of the sick.

*Stimulants.*—The stimulants in common use are *alcohol*, *tea*, *coffee*, and *cocoa*, the last-named deserving to rank among foods. Alcohol is a typical stimulant; it acts as a whip, causing a temporary acceleration of physiological activity. Such acceleration must subsequently be paid for, the extra expenditure brought about by alcohol entailing diminished capacity for further exertion. Alcohol is thus of service only for emergencies of *short* duration; it is eminently harmful when *prolonged* exertion and endurance are required. Like all rapid stimulants, alcohol is in large doses a direct depressant. Tea and coffee owe their stimulating property to the alkaloid *caffein*. They are more useful than alcohol because less liable to abuse, and less dangerous when taken in excess. Cocoa is a stimulant by virtue of *theobromine*, a food by virtue of the large amount of fat or oil that it contains.

The consideration of cost is of great practical importance and not without physiological interest in questions of diet; for the supply-price is a determining factor in the diet-scales unconsciously selected by large masses of men, and laid down by the governing authorities of prisons and of charitable institutions.

The points that the table given below teaches or illustrates are: (1) That nitrogen is much more costly than carbon. (2) That carbon in fat is more costly than carbon in starch or sugar.

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centimeters, and then proceeding to take percentages. It is easier to estimate diets by elements than by proximate principles, heed being taken that carbon has a higher value in fat than in carbohydrate. The real value of a diet often falls short of its paper value.



(3) That nitrogen in an animal proteid is more costly than nitrogen in a vegetable proteid.

The table also accounts for the fact that the poorer classes obtain their carbon by sugar rather than by butter, and that, even in the country, milk and eggs rank as luxuries. Beer is recognised to be an extravagance; rice, which is a staple food among Oriental nations, and peas, which are a representative article in a vegetarian diet, are particularly economical. And in the prices of the three articles, bread, oatmeal, and potatoes, which are most largely consumed by the masses in England, Scotland, and Ireland respectively, we can recognise that the Scotchman gets most for his money, the Irishman least.

COMPARATIVE COST OF SOME COMMON FOOD-STUFFS.

Food-stuffs	Market prices assumed as the basis of calculation	Number of grammes obtained for one penny	
		of Carbon	of Nitrogen
Milk . . .	2 <i>d.</i> per pint	20	1·7
Beef . . .	10 <i>d.</i> per pound	7	1·5
Bread . . .	1½ <i>d.</i> „	80	3·75
Potatoes . . .	½ <i>d.</i> „	90	2·7
Oatmeal . . .	1¾ <i>d.</i> „	100	5
Dried peas . . .	1¼ <i>d.</i> „	110	12
Butter . . .	12 <i>d.</i> „	27	0·06
Eggs . . .	12 <i>d.</i> per doz.	7·5	1
Rice . . .	1½ <i>d.</i> per pound	120	2·2
Sugar . . .	1½ <i>d.</i> „	120	—
Cheese . . .	6 <i>d.</i> „	26	4
Cocoa powder . . .	10 <i>d.</i> „	25	1
Beer . . .	2 <i>d.</i> per pint	14	0·4

According to the now discarded theory of Liebig, proteids were regarded as plastic, fats and carbohydrates as calorific. Bidder and Schmidt considered that only part of the proteid was plastic, and that the remainder was oxidised wastefully, or, as they termed it, that it underwent *luxus consumption*. These views are abandoned; the only sense in which a *luxus consumption* of proteid can be presumed to occur is by the ‘short-cut’ conversion alluded to on page 249; and we are about to learn that *all* food is calorific. The dominant doctrine since 1873 is that of Voit and of Fick, and we have followed it in the above exposition, more especially in that portion which is presented on pp. 256–260. But it should be stated that its fundamental principles have recently been called in question by Pflüger. Whereas, according to Voit, body-fat is produced from food-proteid, but not from carbohydrate, and according to Fick the work of the body is effected at the expense of non-nitrogenous food, Pflüger maintains that body-fat can be produced by food-carbohydrate, that Voit’s analyses are not valid evidence

of the production of fat from proteid, and that the work of the body is effected mainly at the expense of proteid food. That proteid alone can be the adequate source of heat *plus* work is clearly demonstrated in the case of dogs fed exclusively upon lean meat, but Pflüger goes further, and maintains that proteid is the chief driving food, even when accompanied by other kinds of food.

### ANIMAL HEAT

**Physical data.**—The *thermometer* is an instrument containing a fluid, usually mercury, which expands by heat. The expansion, and consequently the intensity of the heat that affects it, is measured in degrees; these are the conventional parts into which the total expansion between the freezing and boiling points of water is divided. The scales of degrees in common use are those of Fahrenheit and of Celsius. On the Fahrenheit scale the total expansion is divided into 180 parts or degrees, the freezing point is at  $32^{\circ}$ , the boiling point at  $212^{\circ}$ . On the Celsius or centigrade scale the total expansion is divided into  $100^{\circ}$ , the freezing point is at  $0^{\circ}$ , the boiling point at  $100^{\circ}$ .

*The calorimeter.*—*Sources of error.*—*Heat-units.*—Amount of heat is measured by the calorimeter, intensity of heat or temperature by the thermometer. Two vessels of water, containing respectively 1 and 2 litres, may be at the same temperature measured in degrees above zero; the first possesses, however, half as much heat as the second, or if the same two vessels of water possess the same amount of heat, then the water in the first vessel must be at a temperature double that of the second. A man weighing 70 kilos possesses twice as much heat as a boy of 35 kilos, the temperature of both being the same; a fever patient weighing 60 kilos, with a body-temperature of  $42^{\circ}$ , possesses the same amount of heat as a collapsed person weighing 70 kilos with a body-temperature of  $36^{\circ}$ .

The unit of heat is the *calorie*, *i.e.* the amount of heat required to raise 1 gramme or cubic centimeter of water  $1^{\circ}$  centigrade; 10 calories are an amount of heat sufficient to raise 10 grammes of water  $1^{\circ}$ , or 1 gramme of water  $10^{\circ}$ , or 5 grammes of water  $2^{\circ}$ , &c.; thus the amount of water in grammes or cubic centimeters, multiplied by the temperature in degrees, gives the amount of heat in calories. The units of heat further employed to express physiological results are the *kilo-calorie* and the *milli-calorie*, the first being the unit generally used in numbers relating to the total heat-production of animals, the latter that used in delicate measurements made on excised muscles. The kilo-calorie is 1,000 calories, or the amount of heat required to raise one litre or kilo of water  $1^{\circ}$ ; the milli-calorie is  $\frac{1}{1000}$  calorie, or the amount of heat required to raise 1 milligramme of water  $1^{\circ}$ .

The calorimeter, as used in physiology, is essentially a chamber within which an animal can be confined so as to impart all the heat



it produces to a known volume of water contained in a surrounding chamber; an external chamber packed, with non-conducting material, limits as far as possible loss of heat by the water, and the temperature of the latter is read upon thermometers. Let us suppose that the water-chamber contains 10 litres of water, which are raised from  $20^{\circ}$  to  $25^{\circ}$ , viz.  $5^{\circ}$  in one hour, by a cat or a dog confined in the inner chamber. We know therefrom that the cat or dog has given off 50 kilo-calories.

*Sources of error.*—But if the theory is simple, the sources of experimental error are numerous and considerable. It is difficult to determine accurately the mean temperature of a large quantity of water; the material of which the apparatus is constructed absorbs heat; the non-conducting envelope is not a perfect insulator; the

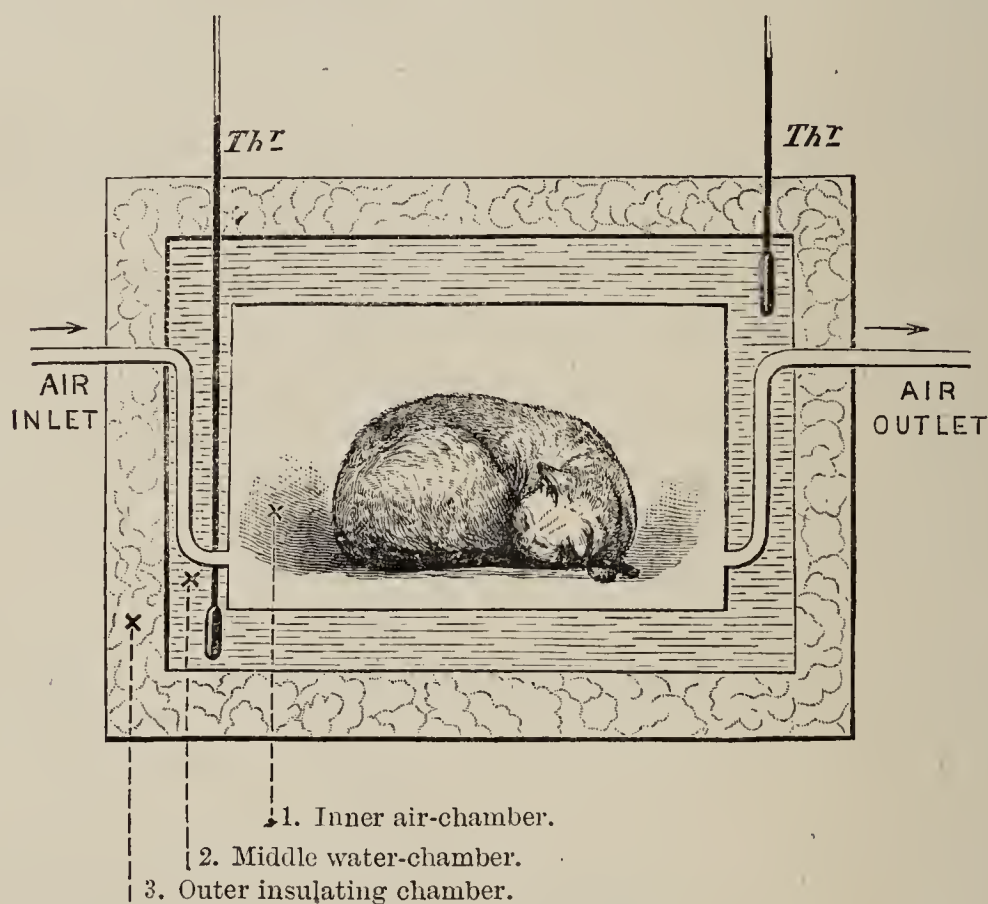


FIG. 111.—CALORIMETER.

temperature of the animal may have risen or fallen; a current of fresh air, which carries off heat and moisture, must be made to enter the chamber. All these circumstances must be taken into account and allowed for as accurately as possible, but it is obvious that a considerable margin of error remains unavoidable.

*Mechanical equivalent of heat.*—Energy cannot be destroyed; it can only be transferred from one place to another. Its various forms are heat, work, chemical action, and electrical action, and it has been experimentally determined how much of one form of energy is equivalent to how much of another form. Thus, it has been determined by Joule that 1 heat-unit is equivalent to 424 work-units. The heat-unit is the calorie; the work-unit is that amount of work required to raise 1 gramme to a height of 1 meter, and is called the grammeter; the

kilo-grammeter and the milli-grammeter are its multiples by 1,000 and by  $\frac{1}{1000}$ . Therefore

1 calorie = 424 grammeters.

1 kilo-calorie = 424 kilo-grammeters.

1 milli-calorie = 424 milli-grammeters or 424 gramme-millimeters.

*Specific heat capacity.*—In the case of animal bodies this may be taken as  $\frac{9}{10}$ ths that of water; *i.e.* an animal requires  $\cdot 9$  kilo-cal. per kilo to raise it  $1^\circ$ , or gives off  $\cdot 9$  kilo-cal. per kilo in falling  $1^\circ$ .

*Latent heat of evaporation.*—To appreciate the relation between evaporation from the skin and consequent heat-dissipation from the body, it is necessary to bear in mind the *latent heat* of water, *i.e.* the amount of heat required to convert 1 c.c. of water into water vapour (or, *vice versa*, the amount of heat set free in the transformation from vapour to liquid). This amount varies with temperature, and may under ordinary physiological conditions be taken as 600 calories. It is the amount of heat removed by 1 c.c. of water from the body and from the air in its evaporation. A day's evaporation from the skin and lungs of 1,200 c.c. would require 720 kilo-calories, most of which is drawn from the body.

*Mass and surface.*—It is important to realise the relation between the weight or mass of a body and the area of its surface. Animal heat is generated by the substance lost from the surface, and in the

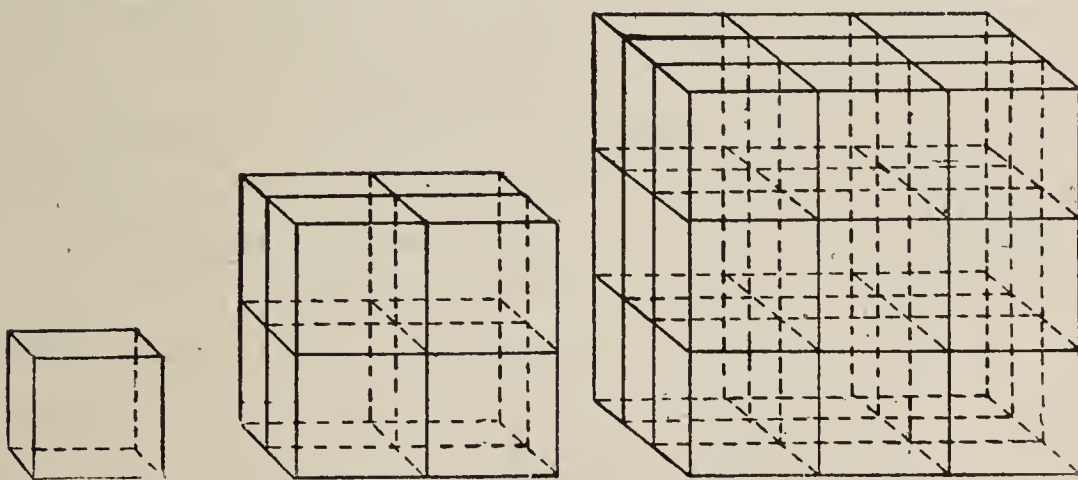


FIG. 112.—DIAGRAM TO ILLUSTRATE THE RELATION BETWEEN VOLUME OR WEIGHT AND SURFACE.

The volumes are	.	.	1	8	27 &c. cubic centimeters.
The weights are	.	.	1	8	27 &c. grammes.
The surfaces are	.	.	6	24	54 &c. square centimeters.
<i>i.e.</i> their ratio of increase is	.	.	1	4	9 &c.

Or otherwise, weight or volume increases as the cube, surface increases as the square; or  $S = 6 W^{\frac{2}{3}}$ .

long run gain and loss must balance. But mass and surface do not vary in proportion with each other; the ratio of mass to surface is an increasing one, as may be easily realised on reference to fig. 112 or to fig. 110, giving the approximate body-surface of a man from 3 to 70 kilos body-weight.

We thus see that in relation to its weight a small animal has a



larger body-surface than a large animal; so that, *cæteris paribus*, a small animal loses heat more rapidly than a large animal; it therefore produces more heat; from which it follows that chemical change must of necessity be more active in a small than in a large animal. The conclusion is, indeed, borne out by experiment; weight for weight, small animals produce and give out more heat, more urea, and more  $\text{CO}_2$ , and consume more food than large animals; and during a given time a larger quantity of blood flows through the tissues of a small than through those of a large animal.

**Animal heat.**—All animals are, as a general rule, warmer than the air or water by which they are surrounded, inasmuch as they are the seat of chemical action. ‘Warm-blooded’ animals (birds, mammals) exhibit this excess of temperature to a far greater degree than ‘cold-blooded’ animals (fishes, reptiles, amphibia), and whereas the temperature of the former varies very little with large variations of the surrounding temperature, that of the latter varies considerably with this circumstance. A man, or dog, or rabbit has a constant temperature, in winter and in summer, at the tropics and in arctic regions, a frog, or fish, has a low or a high temperature according as the surrounding medium is cold or hot. The warm-blooded animal keeps at a constant temperature by reflex actions of a peculiar character, regulating the heat that is produced in the substance of the body, and the heat that is lost from the surface of the body; the cold-blooded animal has no such regulating power, and consequently, no fixed or normal body-temperature.

**Thermotaxis.**—Let us at once look more closely into the manner in which an animal of constant temperature maintains that state. Theoretically, this can be effected in two ways—by variation of the supply, and by variation of the loss—and, as a matter of fact, both these means act concurrently, though the second is the more obviously effective. Heat is supplied by chemical action, and in small proportion by the frictions and concussions that accompany mechanical movements, but these also originate from chemical changes. All chemical actions taking place in the body are originally derived from food, and, as we shall see, the heat-value of the day’s food corresponds with the amount of heat *plus* the amount of work given out. More chemical action, more heat; less chemical action, less heat. It has been experimentally shown that in a medium of low temperature a man gives off more carbonic acid than in a medium of high temperature, and it is a matter of common experience

that he takes violent exercise to warm himself. These facts are sufficient to prove that the supply of heat is diminished in a hot medium, increased in a cold medium. Heat is dissipated by radiation and by evaporation from the surface of the body, only a small fraction (about  $\frac{1}{25}$ th) of the total goes in the discharge of warm air and warm urine; in hot weather the skin is flushed with blood and moist with perspiration, and superfluous clothing is put off; in cold weather the skin is pale and dry, and extra clothing is put on; in the first case the dissipation of heat from the body is accelerated, in the second case it is retarded.

We may in respect of heat-regulation compare the body with a room in which heat is generated by a fire and lost through a window. If it is cold outside the fire burns high and the window closes; if it is warm outside the fire burns low and the window opens, and so the temperature of the room remains unaltered in the two cases. The analogy would be drawn closer, and include the manner of heat-distribution by the circulating blood, if we considered a room or a building warmed by hot-water pipes; the water passing through furnaces representing the blood passing through glands and muscles, and the external wall of the room or building representing the skin of the body; the only feature of difference in the two cases is that the hot fluid system is far more minutely ramified in the body than in a building, forming in the former a cutaneous capillary network which is the chief agent of heat-dissipation, directly by radiation and indirectly by the maintenance of sudorific action. This cutaneous sheet is the true analogue of the window, and is the principal regulator of body-temperature.

We have seen that at high temperature a man makes less heat, is comparatively quiescent, has a flushed and perspiring skin, and clothes himself lightly. At a low temperature he is comparatively active, has a pale, dry skin, and clothes himself thickly. In these two opposite states the heat-production and heat-dissipation concur to produce the required correction by varying in opposite senses, viz. *minus* production with *plus* dissipation, *plus* production with *minus* dissipation. It happens otherwise when increased heat is produced simply and solely as the effect of increased tissue-activity, *e.g.* by muscular exertion; in this case we have increased production with increased dissipation, and as the result only a slight rise of body-temperature. And if heat-production be diminished, its consequence is a diminution of heat-dissipation, which limits loss of heat and



prevents any marked fall of temperature. Thus, in these two opposite states the factors concur to maintain constancy of body-temperature by varying in the same sense, viz. *plus* production with *plus* dissipation, *minus* production with *minus* dissipation.

*Heat-distribution.*—At any given moment the temperature is approximately uniform throughout the body, *i.e.* between  $37^{\circ}$  and  $38^{\circ}$ ; this uniformity is effected by the circulation of blood, which carries heat to parts where heat is lost, and carries off heat from parts where heat is produced; the blood thus warms cooler parts and cools warmer parts. Within the narrow limits of comparative uniformity, slight differences of temperature are detectable in different parts and in their venous blood, according as such parts are giving or taking heat to or from the blood. The mean temperature of the blood is  $38^{\circ}$ ; the warmest organs of the body are the liver, the brain, and the muscles; the coolest parts of the body are the skin and the extremities. The venous blood returning from the intestines during digestion is warmer than the aortic blood, and becomes warmer still in its passage through the liver. The venous blood of an extremity, although coming from muscle as well as from skin, is as a whole cooler than the arterial blood; the venous blood in the *profunda femoris vein*, which is almost exclusively muscular, is a little warmer than arterial blood (about  $\cdot 2^{\circ}$ ). We may illustrate and summarise the thermic relations just discussed in the following figures, selected from various sources, principally from Bernard's observations on dogs, bearing in mind that in this animal the body-temperature is normally somewhat higher than in man.

Average of arterial blood	.	.	$38^{\circ}$ C.	
Blood in left ventricle	.	.	$38\cdot 2$	
Blood in right ventricle	.	.	$38\cdot 4$	
Blood in crural vein	:	.	37	
Blood in profunda femoris vein	.	.	$38\cdot 2$	
Blood in portal vein	.	.	38 to 39	} according to the state of digestion
Blood in hepatic vein	.	.	$38\cdot 4$ to $39\cdot 5$	
Brain	.	.	40	
Liver	.	.	41	
Muscle	.	.	$38\cdot 2$ to 39	(relaxed and contracting)
Skin of trunk	.	.	35 (or lower)	
Skin of extremities	.	.	30	„

These are, however, not to be taken as the absolute temperatures discoverable at any one time in a single animal; practically we are restricted to the comparison between any two simultaneous temperatures, and frequently meet with apparent

incongruities in the course of successive comparisons ; the figures have been selected to exclude these accidental incongruities. And we may call attention to the fact that, although the elevations of temperature are very small, even in the case of glands and muscle, which are the chief heat-producing organs, the absolute heat-production indicated by a small elevation of temperature is very considerable if we take into account the absolute amount of blood passing through the organs ; in the case of the liver, for instance, taking the moderate estimate of 500 litres blood per diem raised  $\frac{1}{2}^{\circ}$ , we have a daily heat-production of 250 kilo-calories, or one-tenth that of the entire body.

*Inflammation.*—Brief allusion may be made to the question of heat-production by *inflamed* parts. The name is suggestive of heat-production, but experiments do not justify us in the belief that more heat is produced in an inflamed part, although its temperature is higher than that of a symmetrically-placed normal part ; the raised temperature is due to vascular dilatation and increased blood-supply, and has never been found to exceed that of arterial blood ; to prove that an inflamed part actually produces more heat than usual, it would be necessary to find a temperature of the part or of its venous blood *above* that of the arterial blood, which has not been done.

*Nerves and nerve-centres concerned in thermotaxis.*—Heat-production, heat-dissipation, and heat-distribution are one and all under control of the nervous system ; they further vary together in such a way that a constant body-temperature is maintained. This harmonious and duly co-ordinated action is what we have referred to above as thermotaxis. Proof that *heat-production* is under the control of the nervous system—the temperature of a limb, paralysed by section of its efferent nerves, falls, chemical change is diminished, and the blood returning through the veins contains less  $\text{CO}_2$  than usual. Proof that *heat-dissipation* is controlled by nerves—if a young animal (kitten) is placed in a warm chamber, the vessels of the skin dilate and the paws begin to sweat, changes that do not occur in a limb the nerves of which have been divided. Proof that *heat-distribution* is controlled by nerves—section of a rabbit's sympathetic causes dilatation of the blood-vessels of the ear and increased temperature ; excitation causes the reverse effects.

These are the simplest among many more or less simple cases in which the production, dissipation, and distribution of heat are influenced through nervous channels. The question



next arising is, whether nerves produce heat-variations by an immediate action upon the tissues, or by the intermediation of vascular changes, or as the concomitant of increased chemical change. It is proved that nerves influence the dissipation and distribution of heat by constricting or dilating the vessels that carry the warm fluid, and by exciting or failing to excite the action of the sweat-glands. It is also proved that nerves influence the production of heat by way of the thermal and chemical effects that accompany activity of muscular and glandular tissues. But as regards the direct action of nerves upon heat-production, apart from the vascular and chemical factors, this has never been demonstrated; thermogenic or thermo-inhibitory action of this character is 'unproven,' and must therefore, in default of evidence, be held to be non-existent. Nor have we any experimental right to assume the existence of thermogenic nerve-centres. Doubtless the nerves that control the three heat-processes, by controlling the calibre of blood-vessels and the activity of muscles and glands, are connected with nerve-centres that act and are acted upon by these nerves, and that therefore influence the heat-processes. Definite experiments prove this fact; excitation and lesion of particular parts of the brain and cord cause the temperature of the body to vary. But the changes of temperature are dependent upon vascular and chemical changes, and we have no more right to assume the existence of specific thermogenic or thermo-inhibitory centres than we have to assume that of thermogenic nerves or nerve-fibres. Indeed, heat-production, unless as the consequence of mechanical or chemical action, is not physically conceivable. This last consideration leads us to recognise that the question of a thermogenic nerve-mechanism is not in reality an independent one, but a part of the much-vexed question of trophic nerves influencing tissue-metabolism, which will be considered in a future chapter.

The allusion above made to the influence of particular parts of the brain and cord on body-temperature is founded upon the following observations:—Clinical cases are not uncommon of injury to the spinal cord, followed by extremely high temperature; others are met with in which the temperature is subnormal. Experimentally, it has been found that punctures of the brain, more especially such as implicate the corpus striatum and optic thalamus, cause a rise of temperature ( $1^{\circ}$  to  $2^{\circ}$ ), which is proved to be due to increased heat-production by direct calorimetry (Richet), as well as by the increased respiratory exchange (Aron-

sohn and Sachs). Obviously, however, these effects are no proof of the existence of direct *thermogenic* nerve-centres.

The thermal effects that accompany the varying activity of glands are experimentally illustrated by the case of the sub-maxillary gland of the dog; the temperature of the venous blood and of the saliva is greater during the activity of the gland than during its rest, and may then exceed that of the carotid blood. The liver affords a still more pronounced instance; the temperature of the hepatic blood is generally higher than that of the portal blood, and the difference is preserved during digestion, when both kinds of blood are raised in temperature.

*Normal heat-production of man. Direct estimation.*—Direct calorimetrical estimates on man give, as the amount of heat daily produced by a man of average weight, 2,500 kilo-calories, or, roughly, 100 kilo-calories per hour. It has been found further that the heat-production is least during sleep, greatest during muscular exertion, being approximately—

during sleep	.	.	.	.	.	40 kilo-calories per hour.
„ rest	.	.	.	.	.	100 „ „
„ moderate movement	.	.	.	.	.	150 „ „
„ active exercise	.	.	.	.	.	250 to 300 „ „

*Indirect estimation.*—The heat-value of a substance is the amount of heat that is produced by its complete oxidation, and this amount is the same whether the oxidation be quick or slow, reached by a direct or by a circuitous path; a gramme of sugar or fat burned in a calorimeter gives the same amount of heat as a gramme of sugar or fat used up by the body. It is therefore possible to estimate the amount of heat that must be produced in the body by estimating the heat-value of the food daily consumed. In the case of *proteid* we must take into account that the excreted urea is an incompletely oxidised product, the heat-value of which must be subtracted from the heat-value of *proteid*.

The heat-value of 1 gramme	proteid	.	.	=	5 k.-cal.
„	fat	.	.	=	9.07 „
„	starch	.	.	=	3.9 „
„	urea	.	.	=	2.2 „

Therefore in a day's diet—

The heat-value of 140 grammes	proteid	.	.	=	700 „
„	105 „ fat	.	.	=	952 „
„	420 „ carbohydrate	.	.	=	1,638 „
„	(-40 „ urea)	.	.	=	(-88) „
	Total	.	.	=	3,202 „

The amount of heat that the above diet is capable of pro-



ducing is about 3,200 kilo-calories. The whole of this amount does not, however, manifest itself as such. The animal body is equivalent to a fuel-fed machine; the *total* energy of the food or fuel becomes kinetic in two forms: (1) work of the body or machine, (2) heating of the body or machine; and in the human body the total energy appears in the following proportions—work  $\frac{1}{5}$ , heat  $\frac{4}{5}$ . The amount of heat actually generated, and sensible as such, is therefore  $\frac{4}{5}$  of 3,200, *i.e.* 2,560 kilo-calories—an amount which is not very different from that obtained by direct calorimetry.

An indirect estimate may also be formed by calculating the heat-values of C and H contained in the two chief oxidation products,  $\text{CO}_2$  and  $\text{H}_2\text{O}$ . Taking as our data an excretion of 852 grammes  $\text{CO}_2$  and 140 grammes  $\text{H}_2\text{O}$  formed in the body, we obtain 1,700 and 500 kilo-calories as the heat-values of C and H in these amounts. But this estimate is of little practical value, owing to the uncertainty of the  $\text{H}_2\text{O}$  estimate taken as a basis; it is only useful to help us to realise the energy-equation of an animal body.

*Thermometry—its scope.*—By the thermometer we ascertain the temperature of the body, not the amount of heat that it produces or loses. The thermometer gives no measure of the activity of heat production, nor even any positive indication that it is above or below the normal. This must obviously be so when we reflect that the amount of heat in a given body, and consequently its temperature, is the resultant of two factors—heat-production and heat-dissipation—either or both of which may vary. Theoretically, of course, temperature varies directly with amount of heat produced, and inversely with amount of heat dissipated, but practically the thermometer does not show whether more or less heat is being produced or dissipated. For instance, in *fever* the body-temperature is usually higher than normal, but we cannot know from this fact alone whether the rise is due to increased production or to diminished dissipation, and it is not surprising that the opinions of authorities have been various on this important point, when reliance has been placed on the thermometer alone.

Enumerating the conditions of which high temperature may be the resultant, we may have any one of the following pairs—normal production with diminished loss; increased production with normal loss; increased production with diminished loss; increased production with increased loss, the former exceeding the latter;

diminished production with diminished loss, the latter exceeding the former. Evidently a rise of temperature is not a simple datum : so far is this from being the case that we had better put into tabular form all the various events that may be concealed under a rise, or fall, or maintenance of body-temperature. And to make the points clearer we have taken imaginary numbers to illustrate variations of gain and loss, putting the normal at 100—which, if so minded, the reader may take to signify calories per hour. The cases in which ‘fever’ would be said to exist are indicated by asterisks.

DATA.		CONSEQUENCE.	
	Heat produced	Heat dissipated	Body-temperature
1	Normal . 100	Normal . 100	Normal
2*	Increased . 150	Increased . 150	Normal
3	Diminished . 50	Diminished . 50	Normal
4	Normal . 100	Diminished . 50	+
5*	Increased . 150	Normal . 100	+
6*	Increased . 150	Diminished . 50	+ +
7*	Increased . 150	Increased . 125	+
8	Diminished . 75	Diminished . 50	+
9	Normal . 100	Increased . 150	—
10	Diminished . 50	Normal . 100	—
11	Diminished . 50	Increased . 150	— —
12	Diminished . 50	Diminished . 75	—
13*	Increased . 125	Increased . 150	—

It is obvious that from simple thermometric observations no conclusion can legitimately be formed. If the essential phenomenon of *fever* be increased heat-production by increased tissue-metabolism, then it is obvious that there may be fever with a body-temperature above, at, or below normal, and as a matter of fact it is admitted by all reliable clinical authorities that the febrile process may be intense when body-temperature is normal or even low.

It would be a valuable supplement to our precise knowledge of fever in its different stages if, failing the calorimeter, which cannot be used clinically, thermometric indications were supplemented by systematic readings of excreted carbon and nitrogen : these give measure of tissue-change, and are obtainable—not, indeed, without great care, but without great practical difficulty.

If the reasoning above given has been properly followed, it is obvious why the important and apparently simple question—‘Is the increased body-temperature of fever the result of increased heat-production, or of diminished heat-dissipation (=heat-retention), or of both factors?’—has not even now, with the aid of calorimetry and thermometry combined, received its final and



assured answer. It is highly probable that both factors come into effect at different periods of a febrile attack.

Considered from the experimental side, the possibilities exhibited in the table show that great caution is necessary in drawing conclusions, even from calorimetrical observations on animals. By the calorimeter we read heat-dissipation, and infer heat-production; a larger calorimeter reading with a *falling* body-temperature is bad evidence of increased heat-production; to be good evidence it must occur with a constant, or, better still, with a rising body-temperature. A smaller calorimeter reading with a rising body-temperature signifies heat-retention; to be good evidence of diminished heat-production it must occur with a constant, or, better, with a falling temperature.

The internal temperature of a bird is high, that of a varnished rabbit falls rapidly, yet the heat-production of a bird is not high, nor is that of the varnished rabbit necessarily low; the high and the low temperatures are respectively surface-effects, heat-dissipation being retarded by feathers, accelerated by varnishing. An infant needs to be kept warm on account of the great heat-dissipation, an aged person on account of the smaller heat-production.

*Surface-temperature and deep temperature:*—It is hardly possible to employ calorimetry for clinical study, but with a due appreciation of the relation that must obtain between temperatures of the surface and of internal parts, it is possible to use the combined data for an approximate estimate of any very considerable alterations of heat-production or of heat-dissipation. To make the matter clear we may consider the two conceivable extremes by which a febrile temperature of the body may be produced—(1) an abnormal heat-production; (2) an abnormal heat-retention. In both cases the internal temperature will be high, but in the first case the surface-temperature will be higher than in the second. The method of taking a surface-temperature is of some importance; it is not possible to obtain an *absolute* surface-temperature, for the instrument applied to the skin protects the subjacent area, and causes its temperature to approximate more and more towards that of deeper parts. Seeing that the information desired is: ‘How quickly is the skin giving off heat?’ and remembering—(1) that in accordance with Newton’s law the rate of heat-emission varies as the *difference* of temperature between surface and surroundings, (2) that heat is also lost by evaporation, we may gauge the heat-dissipation of, *e.g.*

the forearm, from readings of a thermometer and of a hygrometer (w.). The calorimetric value of the thermometer scale is previously determined on a surface giving off heat at a known rate; that of the evaporation is calculated from the weight of water absorbed by the hygrometer.<sup>1</sup> For example: Temperature of air, 20°: thermometer or forearm, 30°; hygrometer covering 20 sq. cm. of skin absorbs 0.5 milligramme H<sub>2</sub>O per minute. (By previous graduation it has been determined that each degree above surrounding temperature indicated by the thermometer strapped to a bag of water cooling at a known rate, signifies a heat-emission of 0.0125 cal. per sq. cm. per min.) The surface of the forearm (not including the hand) is 540 sq. cm.; its heat-emission is then:

$$\left. \begin{array}{l} \text{By radiation and conduction} \quad . \quad 0.0125 \times 10 \times 540 = 67.5 \\ + \text{by evaporation} \quad . \quad . \quad . \quad 0.61 \times 0.5 \times \frac{540}{20} = 8.3 \end{array} \right\} 75.8 \text{ cal. per min.}$$

or, again, *e.g.* after exercise, the thermometer is at 34°, and the hygrometer absorption is 2.5 milligrammes per minute, the emission is then:

$$\left. \begin{array}{l} \text{By radiation and conduction} \quad . \quad 0.0125 \times 14 \times 540 = 94.5 \\ + \text{by evaporation} \quad . \quad . \quad . \quad 0.61 \times 2.5 \times \frac{540}{20} = 41.2 \end{array} \right\} 135.7 \text{ cal. per min.}$$

Simple clinical thermometry, although from a physiological point of view yielding very complex results, is, nevertheless, of

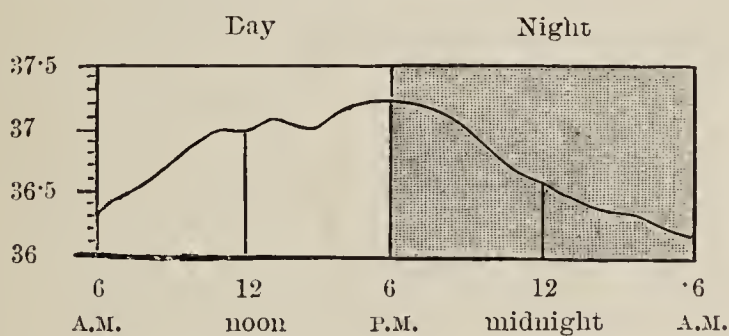


FIG. 113.—NORMAL TEMPERATURE CHART OF THE HUMAN BODY DURING THE TWENTY-FOUR HOURS.

great practical value if carefully performed with reliable instruments, and not committed to the charge of careless assistants. Taken with other indications, the temperature is valuable evidence of the presence and degree of fever and of its probable course. But, to be of use, the observation should be taken twice or four times in the twenty-four hours, and entered for reference upon a temperature chart. The graphic curve of several days is of far more value than a single isolated observation. In view of

<sup>1</sup> P. 246. The evaporation of 1 grm. H<sub>2</sub>O at 30° to 40° requires 610 to 600 calories.



startling statements occasionally made, it is not superfluous to mention that clinical thermometers should possess an index in working order, that the index should be shaken to below the normal point before use, that any suspected instrument should be re-verified, and, finally, that a malingerer may know how to send a thermometer to fever-heat by rubbing it against the blankets. Other memoranda are—the normal daily fluctuations of temperature, the effect of digestion and of exercise, the variations with sex and age. Normally, temperature rises during the day and falls during the night, being thus at its highest at the end of the day, at its lowest at the end of the night. Normally, again, the temperature is raised after a meal and after exertion—more so, indeed, on a weak than on a healthy subject. Finally the temperature is less constant in women than in men, and least constant in children. A high temperature is, as a rule, more significant in a man than in a woman, and in a child it may possibly signify nothing more than a recent fit of crying. The part selected for thermometry may also influence the result; in the axilla the normal reading is  $37^{\circ}$ , in the mouth  $37.2^{\circ}$ , in the rectum  $37.6^{\circ}$ , and the last-named is the most trustworthy.

After an animal has been killed, its body-temperature does not at once begin to fall, like that of an inert mass which has been heated to the same degree. The internal temperature may even rise after death, because with arrest of the circulation and respiration loss of heat has been arrested, or at any rate greatly diminished, while the organs and tissues have still continued to produce heat. This *post-mortem* rise is not solely attributable to rigor mortis, for it makes its appearance sooner; it is a manifestation of the residual metabolism of the tissues and organs of the body, and an expression of the fact that their death does not coincide with that of the organism—the individuals outlast for a time the dissolution of the community. No doubt muscle takes a chief share in such effect, and in so far as it does so its rigor must contribute to delayed cooling; we find, moreover, that in cases where rigidity appears early and strongly (*e.g.* tetanus) the *post-mortem* rise and the delayed fall of temperature are most pronounced. But even in such cases rigor is not the sole factor; it is only one factor in the general tissue-vivacity of which it is a sign. On the other hand, if death supervenes as the termination of chronic disease, if at the death of the body the tissues are moribund, there is no *post-mortem* rise of temperature, and the cooling of the body is but little delayed,

in comparison with that of an inert mass. Deaths from fever afford instructive illustration of the varying thermic effects of residual metabolism; the tissues of a fever patient dying at a high temperature are active, and by continuing active retard the cooling of the body after death; the tissues of a fever patient dying at a low temperature are already moribund, and have but little retarding effect upon the further cooling of the body.



## PART II

### THE PHENOMENA OF EXCITATION

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#### CHAPTER VIII

##### *GENERAL PLAN OF THE NERVOUS SYSTEM*

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299	<i>Why and How ?</i>

PROTOPLASM is excitable. When any part of a lump of protoplasm is excited, the lump moves. When many lumps of protoplasm are gathered into a homogeneous mass, excitations and movements may be transmitted from lump to lump in all directions. With higher organisation of the mass, differences of function and of structure begin to make their appearance. Excitability, while still pervading the whole organism, becomes localised with greater intensity in some parts than in others, along some lines more than along others (sense-organs, nerves, and nerve-centres); in other parts contractility becomes the salient character (muscles). To illustrate this progressive elaboration of a nervous system we may select—(1) an amœba, (2) a jelly-fish, (3) a frog, (4) a man.

An amœba is a simple lump of protoplasm, excitable and contractile in all parts of its substance, and not more so or less so in one part than in another.

The lower surface of the umbrella of a jelly-fish is covered with a layer of contractile protoplasm, which is not far removed from a homogeneous state; an excitation applied to any part causes a contraction, which is transmitted in all directions through the mass; it is a sheet with neuro-muscular properties, a network of rudimentary muscle and nerve.

A frog exhibits a far higher degree of differentiation. Instead of a neuro-muscular cell we have a comparatively complex chain of parts, different in structure and in function, muscle-fibre exhibiting contractility as its salient characteristic, nerve-fibre and nerve-cell possessing conductivity without contractility. In such an animal we recognise the component parts of a nervous system as central, peripheral, and intermediate, and, as we shall see, we may further analyse these components into cerebral and spinal centres, into afferent and efferent conductors, and into many kinds of end organs in connection with these conductors.

The progressive elaboration of a nervous system with subdivision of function has its highest expression in the human brain. There is in the human brain, and in less degree as we go down the scale, in the mammalian brain, a localisation of different functions in different parts. We shall find that this localisation is best known in the cases of language, of voluntary motion, and of sight, and that as regards voluntary motion in particular, it is possible to trace a correspondence between particular movements and the functional activity of particular areas of the cerebral cortex.

The literal meaning of the word excitation is 'call from without.' The surroundings of an organism '*ex-cite*' its specially excitable parts, and the organism moves to or from its surroundings, or registers an impression which will modify its future movements. The specially excitable parts are on the external surface, exposed to excitation; they have no direct communication with the specially contractile parts—the muscles, but an indirect communication by strands of specially excitable protoplasm—nerve-fibres, which conduct excitations to central organs, where they are received, while other nerve-fibres establish the communication from the nerve-centres to the muscles, conducting from the former to the latter the impulses that give rise to movements.

The central nervous cell is usually considered as a double cell, one part of which is at the central end of an afferent fibre, the other half at the central end of an efferent fibre. The first is called a *sensory* cell, the second is called a *motor* cell; and we must imagine that a bond of union (commissural fibre or fibrils) establishes communication from 'sensory' to motor cell (but see p. 551).

The component parts just enumerated form a reflex arc, and



may be classified as central, peripheral, and intermediate. The central part is a nerve-centre or centres, the peripheral parts are the sensificatory organs and the muscles, the intermediate parts are the nerves—afferent and efferent—and the fundamental nervous act is a reflex act. In relation to the conduct of the animal body, the sense-organs are the intelligence department, nerve-centres are the headquarters, muscles are the executive, nerves are the channels of communication.

In the ascending scale of organisation there is increasing diversity of parts, increasing subdivision of properties and of functions, with further diversities between the parts of parts. The simple nervous system below-figured is not that of a mammalian animal nor of man. The ‘nerve-centre’ of mammalia and of man is a collection of nerve-centres occupying the cerebro-spinal axis, with more or less diverse special offices under their

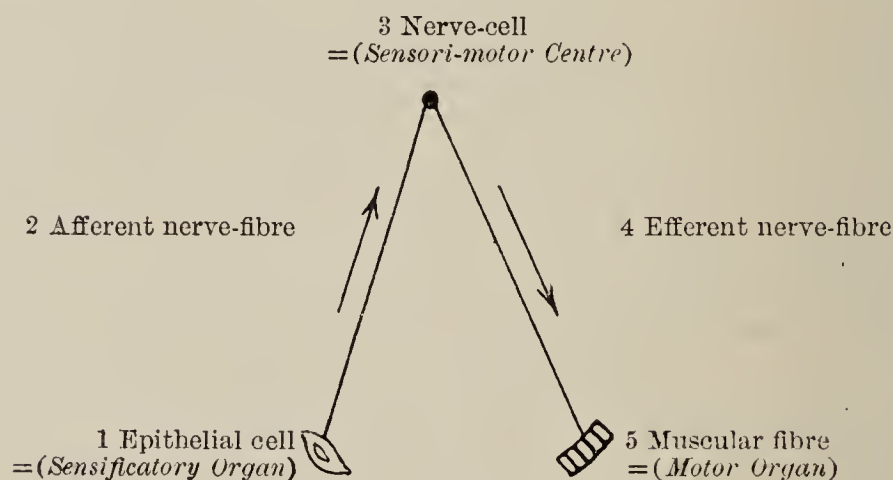


FIG. 114.

control—communicating each with the other upon occasions, yet separately active upon other occasions—having functions that are localised at certain parts, yet not strictly confined to these parts—playing upon and influencing each other in all directions, yet in some directions rather than in others, and maintaining some kind of precedence and rank, so that while all may influence all, yet some are usually guided and controlled by others—variously organised through past excitations, yet still variously organisable by excitations to come. To-day the state and disposition of organs and of the organism are the product of the past, immediate and remote, individual and ancestral. Tomorrow and in the distant future they will become what they may be made to become by training, by education, and by new conditions of life.

**Classification of nerve-centres.**—The cerebro-spinal centres may for convenience be classified as : (1) cerebral or cortical ;

(2) spinal or medullary ; and the reactions of these two classes may be characterised as (1) voluntary ; (2) reflex.

Intermediate between these two classes, partly including both, and absolutely distinguishable from neither, a third class of central action may be recognised—characterised as *automatic* action ; in view of the confusion of thought that has often arisen from the use of this word, it would perhaps be better to avoid it altogether ; at any rate we must define the sense in which it is to be employed. The word has received two diametrically opposed meanings, viz. (1) self-moving, self-arising, spontaneous, in literal translation of *αὐτό-ματος* ; (2) automaton-like, that is to say, like a mechanism that appears to be self-moving, but that we know to be moved by secret springs and hidden keys. The second sense is that in which the word is used in this book. But when we attempt to distinguish the characteristics of an automatic from those of a voluntary or of a reflex action, we are at a loss to find any sharply defining form of words ; at best we may say that an automatic action is one to which at first sight we should be tempted to apply the qualification ‘voluntary,’ but of which some exciting cause, although not immediately obvious, can in general be discovered : usually, but not always, an automatic, like a reflex act, is unconsciously performed. In fact, the automatic action is essentially a reflex action, and differs from it only in that it is, as a rule, the habitual or serial effect of habitual or serial stimuli ; *e.g.* walking, mastication, respiration, &c. Although the word is in certain respects a convenient one to use, we do not see reason for applying it as a classifying term on the same footing as the terms *reflex* and *voluntary*, and shall employ it only as a subordinate term for certain habitual actions belonging to the voluntary or to the reflex categories.

An automatic act is the repeated or rhythmic motor response to a repeated or continuous excitation. Usually it is carried on unconsciously, but it may upon occasion be attended to and its effects felt, and it may be accelerated or restrained by voluntary action. The movements of respiration, of sucking, and of walking, are the most typical among automatic actions ; each of these actions is a more or less prolonged and rhythmic series of motor responses to prolonged or repeated excitation. In the case of breathing, the rhythmic series of movements lasts through life, its chief excitant being the state of the blood and the state of the lungs acting upon the spinal bulb. The automatic act of walking is carried on in the form of more or less prolonged series of movements subject to voluntary control, consciously or unconsciously, but generally unconsciously, and in response to the series of excitations caused by contact with the ground and state of muscular tension. Skilled movements, such as writing or speaking, are on the borderland between automatic and voluntary ; they began as voluntary, and they have later become automatic : even in its fully developed form



each such automatic series of movements is initiated by voluntary effort, and in its highest manifestations is effected in largest measure with the participation of voluntary cerebral action.

**Reflex Action.**—We may define a *reflex act* as the *immediate* motor response to centripetal excitation; it is unchosen, ‘fatal.’ It is subserved by:—

- |   |                                     |
|---|-------------------------------------|
| 1. Epithelium (peripheral sense-organ). | 4. Commissural fibre.               |
| 2. Afferent nerve-fibre.                | 5. Spinal cell on efferent tract.   |
| 3. Spinal cell upon afferent tract.     | 6. Efferent nerve-fibre.            |
|   | 7. Muscle (peripheral motor-organ). |

If the peripheral excitation is weak, a reflex act may be inhibited. If the peripheral excitation is strong, the act may be performed

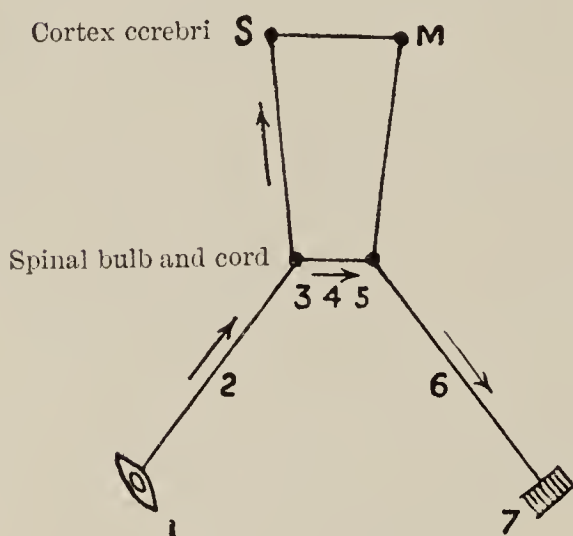


FIG. 115.—REFLEX ACTION.

in spite of an inhibitory effort. It may be performed unconsciously, and never be known to its performer, or it may be unconsciously performed, and become known subsequently, or it may be perceived during its performance.

These several modes and possibilities will be best appreciated by considering examples of each as they occur in everyday life.

The most undeniable of reflex

acts are those of a paraplegic patient or of a headless frog, when in response to cutaneous stimulation the limbs move. The act of winking in response to an irritation of the conjunctiva, or to retinal stimulation, caused by the sudden approach of a missile, is likewise a reflex act. By a voluntary effort the act of winking may be restrained—inhibited; but it may happen that the irritation becomes so irresistible, or that a threatening blow is so sudden, that no voluntary effort can restrain the act; it is a familiar experience with most people that the eyes are continually winking without consciousness having been aroused, and that by paying attention to the matter they may become conscious of the act. Sneezing and coughing, although more complicated, are to be classed among typical reflex actions; a person may sneeze or cough unconsciously or consciously; he may or he may not be able to restrain a sneeze or cough; he may have sneezed or coughed

unconsciously at one time, and yet may subsequently remember the fact, and may then become conscious of having sneezed or coughed.

All the foregoing instances are typical reflex actions, in the performance of which the spinal bulb and spinal cord are the organs of 'return of action.' But the term 'reflex action' has spread beyond these its original limits, and as commonly used at the present day embraces on the one hand visceral and glandular effects, in the production of which the spinal cord is the centre of return of action; and on the other hand includes many highly complex series of movements, in which the bulb or even the cerebral cortex takes a leading part, but which may nevertheless be carried on with or without the participation of consciousness. Instances of the reflex action of visceral muscle are furnished by the modifications of the heart's beat, and of the state of the blood-vessels, that are brought about by centripetal stimuli acting through the spinal bulb and cord, and giving rise to centrifugal impulses along vaso-augmentor or vaso-inhibitory nerves. Of this class of reflex effects, blushing or pallor, fainting or palpitation, are the most familiar instances. It is to be noted that in this, more than in any other class, the reflex effect can most obviously be of two kinds, a reflex increase of action or a reflex diminution of action. Reflex contraction of smooth muscle is also exemplified in the third stage of deglutition, in the act of defæcation, in the case of the urinary and biliary bladders, and in the alterations of the iris consequent upon optical stimuli. Reflex glandular action is well characterised in the case of the salivary and lachrymal glands by the effects that are produced by excitation of the mucous membrane of the mouth or of the orbit.

Turning to the consideration of reflex acts in which the bulb and brain are undoubtedly organs of 'return of action,' we recognise that we have reached to, if not beyond, the furthest limit of the term reflex. Walking, breathing, sucking, skilled movements such as those of an accomplished dancer or musician, acts unconsciously performed but bearing all the marks of intelligent purpose, the unconscious cerebration of a mathematician or of a poet—all these are not unfrequently called reflex and alluded to as reflex actions; 'automatic,' in the sense of 'autmaton-like,' as defined above, is the more correct qualification to use, although if we bear in mind that the term 'automatic' is in reality covered by the term reflex (an automatic act being an habitual or serial reflex), it is not absolutely incorrect to call the actions reflex.



But it is better to distinguish the group of automatic actions out of the larger class of reflex actions, and to refer to them under that designation. These automatic actions are distinguishable into two sub-groups—(1) the primary or inherited automatic acts already considered, (2) the secondary or acquired automatic acts. Respiration is a primary automatic act; writing is a secondary automatic act. Walking occupies a position midway between the primary and secondary group; it is primary in so far as the disposition to walk is inherited; it is secondary in so far as a child must learn to walk by practice.

**Voluntary action.**—To any one who will reflect upon the three words, reflex, automatic, voluntary, it will be evident that the actions so qualified overlap, and that the words themselves can only be arbitrarily defined. We have recognised automatic to belong to reflex action; we shall find that, subjectively viewed, *i.e.* by appealing to our own self-consciousness, we all believe our voluntary actions to be spontaneous, and freely chosen in spite of exciting impulses; objectively viewed in the conduct of living beings as it unfolds itself before us, voluntary action appears as a highly disguised and complicated form of reflex action, with its causal excitations more or less concealed, more or less deeply buried in the past history of the individual, or of the ancestors. From either point of view, it is most difficult to settle upon a satisfactory *differentia* by which to characterise and identify a voluntary action. But this is not to say that voluntary is to be entirely undistinguished from reflex action. On the contrary, the difficulty, having been clearly realised, may be, if not logically surmounted, practically turned, by careful description of typical reflex and of typical voluntary actions. As voluntary actions of the highest order we may instance the working out of a mathematical problem, the opinion uttered by counsel or by physician, the composition of a picture or of an opera, the directions pronounced by a responsible leader. From actions of this order, down to the aimless saunterings of an unemployed person, bent in this or that direction by each trivial incident, the movements of apparently free agents are termed voluntary. No line of demarcation can be drawn anywhere between the more or less voluntary actions included between these two extremes. But on review of the whole scale we recognise that an act is voluntary and cerebral in proportion with the imagination and judgment that intervene between sensation and motion. A single voluntary contraction, a shorter or longer

series of voluntary contractions, the lifelong pursuit of a policy, alike imply the following elements—(1) the consciousness of a desire, *i.e.* sensation; (2) comparison and deliberation of cerebral ideas, *i.e.* judgment; (3) executive action, *i.e.* voluntary motion.

Nor is it possible to lay down at any point a boundary-line between the highest voluntary and the lowest reflex action; the quality of centre, whether of brain or spinal cord, is the resultant of past impressions—individual and ancestral, and upon that quality depends the motor result of each given centripetal impulse and sensation. Contrasting the two extremes, we have in the purest reflex action an immediate, ‘fatal,’ unchosen response to peripheral stimulation; the nervous path is deeply canalised; alternative reactions are few; the particular reaction to a particular stimulus is easily predicted. In the purest voluntary action we have a delayed, ‘free,’ chosen response to a given sensation; the nervous path is not performed; alternative reactions are numerous; the particular reaction to a particular sensation cannot be predicted. Between these extremes all imaginable movements and actions of men and of animals find their place.

From the objective point of view of the observer and experimenter, the most definite *differentia* between reflex, automatic, and voluntary actions is afforded by the time factor. A reflex act is a single *immediate* motor reaction; an automatic action is a *series* of immediate motor reactions; voluntary action is *delayed* motor reaction.

If the doctrine of spontaneous volition be accepted (an admission which seems to entail acceptance of the view that effects may occur without causes, or phenomena without generators), the voluntary act commences at the cortical motor cell. But it is more logical to admit that previous sensations have been registered, and that volition is a resultant of past as well as of present sensations. The cerebral sequence is to feel, to judge, to will, and the principal characteristic of cerebral action is great *delay* of reaction, in contrast with the immediate response characteristic of reflex acts.

A voluntary act is subserved by—

- |                           |                           |
|---------------------------|---------------------------|
| 1. Epithelium.            | 7. Cortical motor cell.   |
| 2. Afferent nerve-fibre.  | 8. Efferent tract.        |
| 3. Spinal sensory cell.   | 9. Spinal motor cell.     |
| 4. Afferent tract.        | 10. Efferent nerve-fibre. |
| 5. Cortical sensory cell. | 11. Muscle.               |
| 6. Commissural fibre.     |                           |



A cerebral act is usually attended by consciousness, but undoubted cerebral acts may be unconsciously performed. A cerebral act is usually voluntary, *i.e.* chosen, but it may be automatic

or reflex, *i.e.* an immediate and involuntary response to peripheral excitation, either as a single action or as a prolonged series of acts.

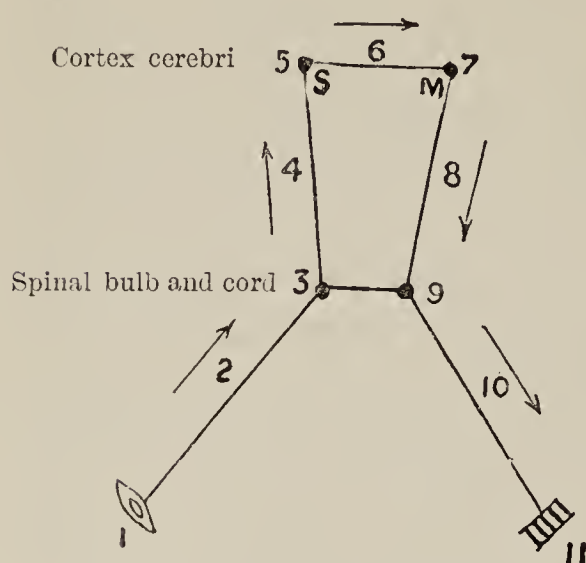


FIG. 116.—VOLUNTARY ACTION.

A familiar comparison may serve to bring home to some minds a clearer picture of the relation that subsists between cerebral and spinal shares in the government of the body, and lead to a rational conception of the relative significance of the terms voluntary and reflex. An ordinary business

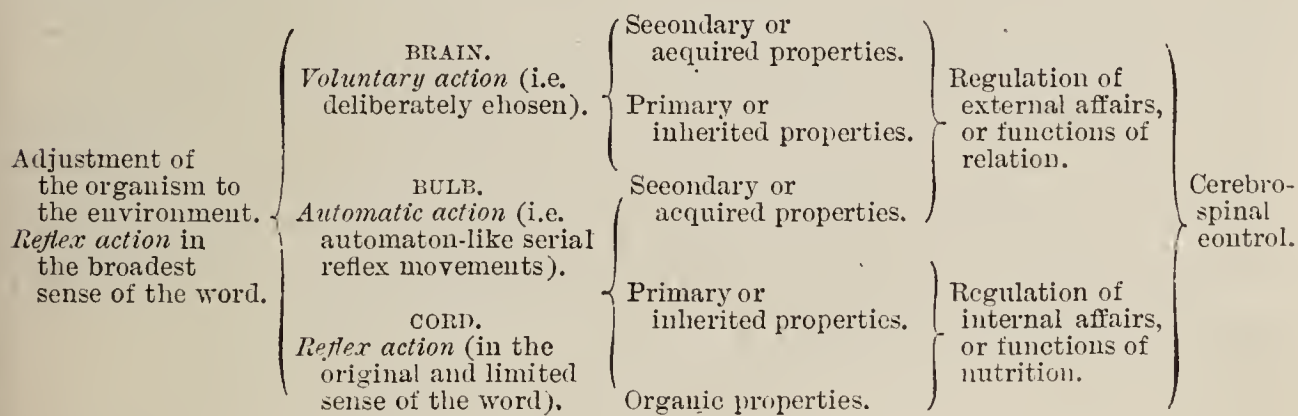
enterprise, a factory, a political party, an army—in short, any body of men gathered together under leadership—is, like the collection of cells forming the animal body, led and controlled by certain individuals whom we may call nerve-centres, and characterise as ‘master centres’ and as ‘foreman centres.’ The function of a master centre in a body, as in a business, is mainly that of administration, to initiate proceedings the detailed supervision of which is delegated to and carried out by foreman centres, to actually take part in supervision only of novel proceedings or of proceedings that go awry; the function of a foreman centre is mainly that of immediate supervision, to execute instructions received from the master centre, to issue instructions in matters of routine without reference to headquarters, but to inform and take instructions from the master centre as emergence arises. This is a true picture of the relations between the cortex cerebri and the medulla spinalis, as far as we know or can conceive them.

The analogy might easily be pushed further, but analogies are to be used with caution; it is, however, allowable to point out (1) that while the mandatory and inhibitory intervention of superior centres is necessary at times, attempts at detailed interference with reactions that are normally within the province of subordinate centres would embarrass smooth working. Thus, a man who should attempt to walk or write by a series of voluntary movements would walk badly or write badly. (2) That although most experimental evidence is to the effect that

those functions which we may regard as the domestic concerns of the body—circulation, respiration, secretion—are regulated by subordinate centres, by the grey matter of the spinal bulb and cord, yet we must recognise from pathological evidence, and from some few physiological experiments, that the cerebrum may, upon occasion, take cognisance of the domestic state as well as take part in its regulation or in its disarrangement.

We may, in conclusion, take this opportunity of pointing out that the relative importance of cerebral and spinal control varies with rank in the animal scale. As we ascend in the scale the importance of the cerebrum increases, that of the spinal axis decreases. The importance of cerebral function is greater, that of spinal function is smaller, in a dog than in a frog, in a man than in a dog. The range and variety of cerebral action increase, whereas the range and variety of spinal action decrease as we ascend in the animal scale. Although the entire cerebro-spinal conduct is more fixed and stereotyped in a frog than in a dog, in a dog than in a man, the increased freedom and variety do not include spinal but only cerebral action ; spinal reactions are more limited and stereotyped in man than in the dog, in the dog than in the frog. The spinal cord is pre-eminently the conservative organ, the cerebrum is pre-eminently the progressive organ.

The considerations conveyed in the preceding pages may now be summarised as follows :



It is necessary, in order to define the scope of the foregoing analysis, to add a few words of limitation. We have followed an idea to the borderland of physical phenomena, which can be measured by our senses. We have glanced across the limit into regions where questions and opinions cannot be measured by objective means, although they have been fearlessly broached and freely pronounced upon by schoolmen and by churchmen. We desire not to trespass upon those regions. The question *why?* is not answered by positive science, but only the question



*how?* and sometimes the question *how much?* Physiological problems are limited to the tangible and measurable phenomena of living bodies. The physiologist cannot say why a muscle contracts, nor define 'life,' 'free will,' 'moral responsibility'; his professed task is limited to an objective study of the essential particulars in which living differs from inert or dead matter; he can learn 'how' a muscle contracts, under what conditions, how much, and with what effects; the original cause of the property termed contractility is beyond his knowledge.

The distinction just drawn is of fundamental importance; it should be emphasised and clearly apprehended that the choice open to us often lies between these two paths: between a vain attempt to answer the question *why?*, to discover a final cause 'a quo' or a final cause 'ad quem': and the feasible task of studying relations between phenomena, of retracing successive steps of '*how*' in the procession of events. We may be able to see how things have happened or will happen, we have no right to say why they have happened, or why they will happen; and it is a first step in the acquisition of positive knowledge to know that the '*ratio rei*' is not the 'reason why.'

In the review of the nervous system upon which we are entering, we shall describe the physiological properties or *modus vivendi* (1) of the peripheral organs—muscle and nerve—(2) of the sense-organs—eye, ear, &c.—(3) of the central organs—spinal cord, spinal bulb, and brain. Animal electricity, although a general phenomenon of chemical change, will, on account of its close association with excitation effects, be considered in connection with muscle and nerve; on the other hand, several subjects belonging properly to the nervous system, *e.g.* the larynx, vascular nerves, &c., have already, of necessity or for convenience, received attention in Part I.

## CHAPTER IX

## MUSCLE

- |      |  |
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THE peripheral nervous system is composed of nerve-fibres and their terminations, and also includes a large proportion of nerve-cells, which enter into the constitution of ganglia and ganglionic plexuses. The peripheral terminations of nerves fall into two chief classes, sensory and motor; the sensory being mainly distributed on the external surface of the body, in the skin and sense-organs; the motor, in the voluntary skeletal muscles. Muscles



may, indeed, properly be regarded as the end organs of motor nerves, and it is in many respects convenient to consider the physiology of muscle in conjunction with that of nerve. Their experimental study necessitates the use of certain electrical instruments and a knowledge of the physical principles involved. These data are therefore presented in their natural place as the physical preface to the physiology of the nervous system.

### ELECTRO-PHYSIOLOGICAL INSTRUMENTS AND PRINCIPLES <sup>1</sup>

The **Daniell cell** is the type of all other cells in common use; it consists of two elements, zinc and copper, and of two fluids,  $H_2SO_4$  and  $CuSO_4$ , separated by a porous septum. The positive element, at which most chemical action takes place, is the zinc, which dips in the sulphuric acid contained in the porous pot; the negative element is the copper, which dips in the copper sulphate solution. The free end

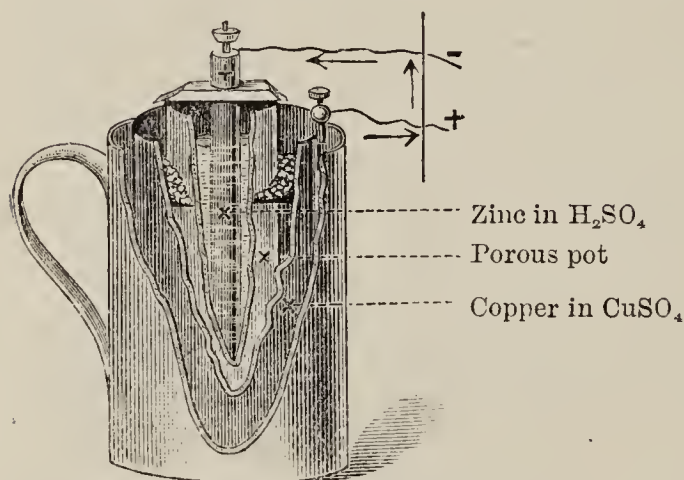


FIG. 117.—DANIELL CELL REPRESENTED SO AS TO EXHIBIT ITS CONSTITUENT PARTS.

of the wire connected with the zinc is the *negative electrode* or *kathode*; that connected with the copper is the *positive electrode* or *anode*. The copper sulphate solution should be saturated, and kept so during use, by an excess of the salt; the zinc should be amalgamated before use, in order to prevent its waste by local action; and when a Daniell cell is required to supply a current of constant

strength, it is well to substitute a saturated solution of zinc sulphate for the sulphuric acid in the porous pot. The direction of current throughout any circuit is from positive to negative—from higher level to lower level; *e.g.* in a circuit composed of a Daniell cell and of wires leading to a nerve laid across electrodes, the direction of current in the cell is from zinc, the positive element, to copper, the negative element; outside the cell, it is from the copper to the anode, through the nerve from the anode to the kathode, and from the kathode to the zinc. As regards the nerve, the anode, or positive electrode, is the seat of entrance of current; the kathode, or negative electrode, is its seat of exit.

**Keys and commutators** of various patterns are used for conveniently making and breaking a current, and for reversing its direction. A key

<sup>1</sup> This section is intended for *practical* students; it may be referred to by other readers for explanation of apparatus alluded to in the text.

in common use is the *Du Bois-Reymond* key, which consists of two metal blocks, each carrying two binding-screws, and joinable at will by a third metal piece that is raised or lowered by a handle. This key can be used in either of two ways : (1) the key can be interposed in one wire of a cell, in which case the circuit is *made* by lowering the junction piece, and *broken* by raising it ; (2) both wires can be brought to each side of the key, from which two other wires lead off to the electrodes ; in this case the circuit through the electrodes is *broken* by lowering the junction piece, and *made* by raising it.

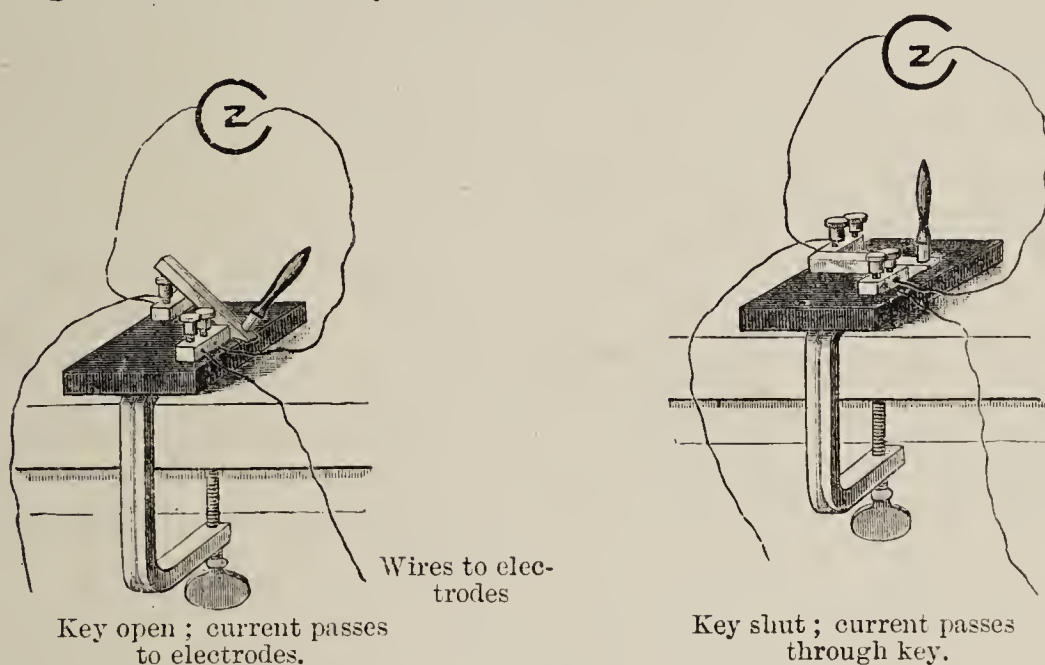


FIG. 118.—DU BOIS-REYMOND'S FRICTION KEY.

Put up so as to 'short-circuit' current when it is closed.

The commutator generally used in a physiological laboratory is *Pohl's mercury commutator*, the construction of which is as follows. A block of insulating material (ebonite or paraffin) is provided with six pits containing mercury, with which six binding-screws fixed in the block

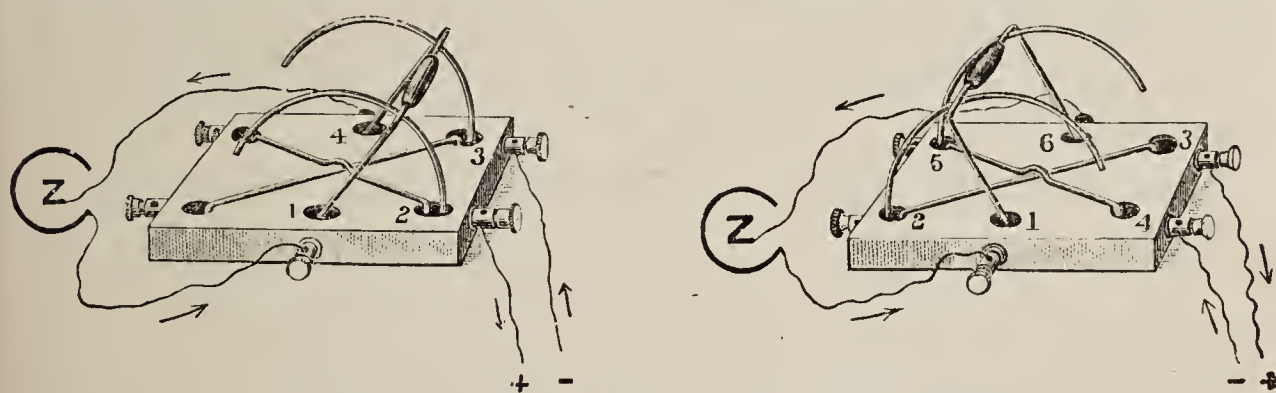


FIG. 119.—POHL'S COMMUTATOR (WITH CROSS-WIRES).

The Daniell cell and electrode wires remaining fixed, the direction of current in the latter is reversed by moving the cradle to right or left. Numbers 1, 2, 3, &c. indicate the path of current in the two cases.

are in communication. Two cross-wires establish junction between the pools of mercury, as shown in the figure. A cradle, consisting of an insulating handle fixed to two metal arcs, can be tilted so that each of the two middle pools is at will brought into communication with



either of the two lateral pairs of pools, right or left. Wires are connected with the middle pair of pools and with one of the lateral pairs. Tilting the cradle to right or left reverses current through an electrode circuit, as can be seen by tracing its path in the figure, or, better, by putting up a circuit consisting of a Daniell cell, key, commutator, and galvanoscope.

A commutator is often used for a different purpose without the cross-wires. In this case both battery wires are connected with the middle pools, two other pairs of wires connected with the two lateral pairs of pools leading off to two pairs of electrodes, which can be applied to two portions of a nerve, or to a nerve and to a muscle. By tilting the cradle to right or left the battery current can now be turned on to one or other pair of electrodes (see fig. 138).

A comprehension of the meaning of *Ohm's law*, and of the terms *current*, *resistance*, *electromotive force*, *potential*, is an indispensable preliminary to the study of electro-physiology. These terms can be sufficiently explained by comparing an electric current with a current of water, although the analogy is in some respects defective. An electric current from a Daniell cell flows through a wire from positive to negative potential, just as the water from a raised reservoir flows through a pipe from higher to lower level. Potential remaining constant, more current flows through a thick than through a thin wire, just as, water-pressure remaining constant, more water flows through a broad than through a narrow pipe. A thin wire offers a greater resistance to the passage of current than a thick wire, just as a narrow pipe may be considered to offer greater resistance than a broad pipe. With a given wire there is more current from higher than from lower potential, just as with a given pipe there is more current from a higher than from a lower water-level. The driving-force is called electromotive force, electromotive difference of potential, potential, or pressure; these are synonymous terms, and the last—viz. pressure—brings out clearly enough the analogy of electrical potential or pressure with water-level or pressure.

**Ohm's law** is simply a short formula embodying the above considerations, viz.  $C = \frac{E}{R}$ , or amount of Current varies directly as Electromotive pressure, and inversely as Resistance. We are already familiar with an analogous but less simple and precise relation in the physics of blood-flow, to the effect that amount of flow increases with increased heart's force, diminishes with increased peripheral resistance. An electrical current can be increased by increasing the electrical pressure, or by diminishing the resistance; it can be diminished by diminishing the pressure, or by increasing the resistance. The principle of electrical measuring instruments may also be understood by comparing them with instruments already familiar to the student of physiology. Amount of blood-flow is ascertained by calculation from the indications

of a dromometer. Amount of electrical flow is ascertained from the indications of a **galvanometer**. Blood-pressure is measured by a manometer; potential or electrical pressure is measured by an **electrometer**, which is practically an electrical manometer.

The conventional names given to the electrical units we employ in physiology are as follows:—

The unit of current is called an **ampère**.

The unit of resistance is called an **ohm**.

The unit of pressure is called a **volt**.

The potential or pressure of a Daniell cell is nearly 1 volt, the internal resistance of an ordinary cell ranges from  $\frac{1}{2}$  to 10 ohms. 1 volt *pressure* through 1 ohm *resistance* gives 1 ampère *current*; through 10 ohms it gives  $\frac{1}{10}$  ampère, &c.

A **rheostat** is a set of resistance coils graduated in ohms, by means of which more or less resistance can at will be put into a circuit.

A **rheochord** is a wire, *r r r*, and slider, *s s*, so disposed that a very low but variable resistance can be offered as a *deriving*, or 'shunt' circuit, by the side of a *principal* circuit of higher resistance. It affords means of dividing a current into two parts, and of thus obtaining any desired small fraction of the current of a single Daniell cell, the larger part passing through the rheochord,

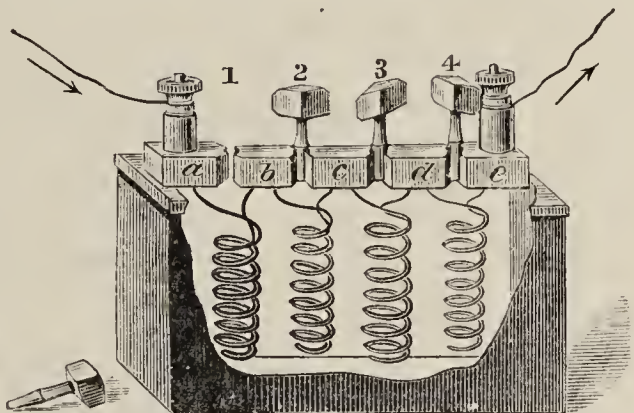


FIG. 120.—RHEOSTAT.

Wires of definite resistance unite the metal blocks *a, b, c, d, e*, which can be connected and disconnected by inserting or removing metal plugs. If all the plugs are inserted, the resistance is practically zero; if a plug is removed (as shown in figure), resistance in a circuit is increased; if the four wires in the box have resistances of 1, 2, 3, and 4 ohms, removal of all the plugs would give a resistance of 10 ohms.

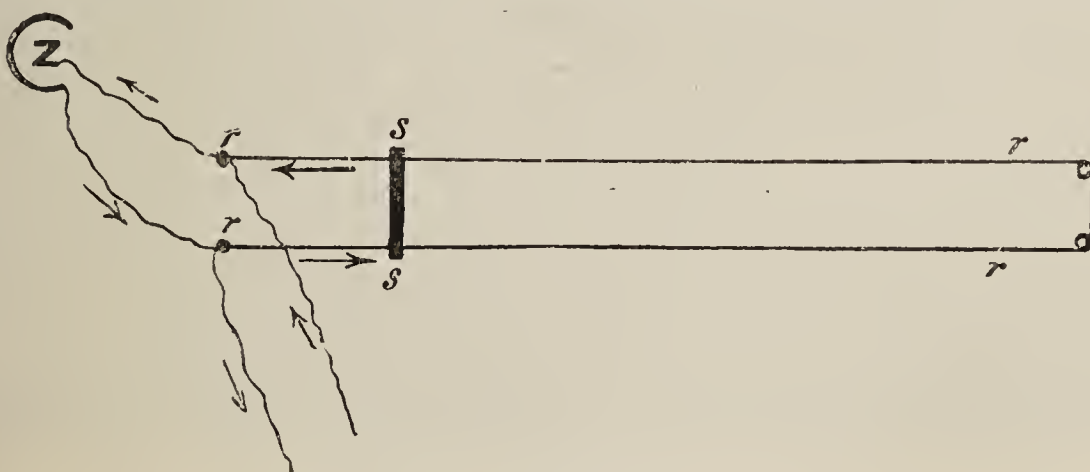


FIG. 121.—A SIMPLE RHEOCHORD.

A key (not shown in figure) is to be used between the battery and the rheochord, not between the rheochord and electrodes.

*r r r*, which is of small resistance, the smaller part passing through the principal circuit, which is of high resistance, and includes the



electrodes and the nerve or muscle. If, for instance, the current of a Daniell is made to branch (a) through a rheochord with a resistance of 1 ohm, (b) through a nerve with a resistance of 9,999 ohms, then the current in the shunting or rheochord circuit will be  $\frac{9999}{10000}$ , the current in the nerve circuit will be only  $\frac{1}{10000}$  of the entire current of the Daniell cell. The rheochord as a deriving circuit is to the principal

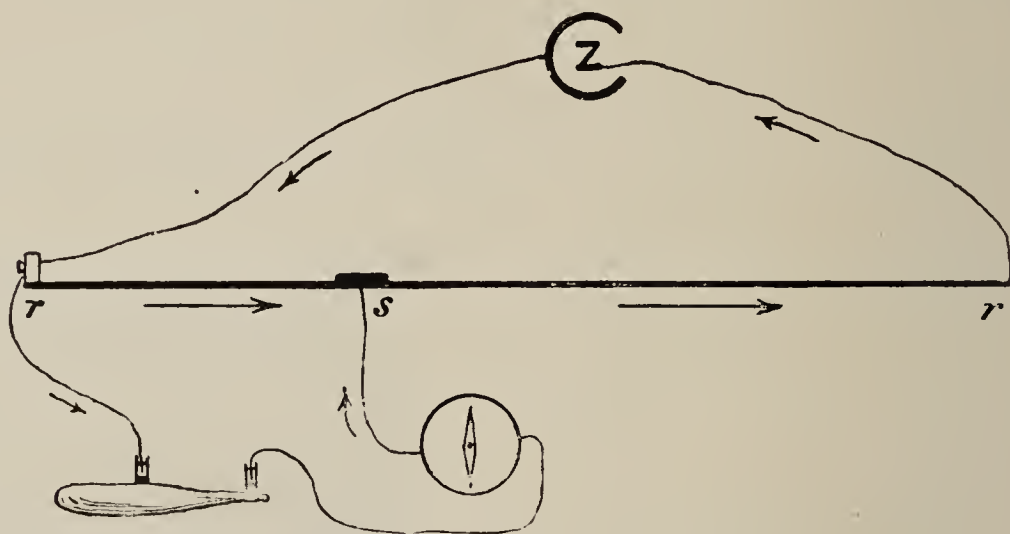


FIG. 122.—A STRAIGHT RHEOCHORD (MONOCHORD) AS DISPOSED FOR COMPENSATION.

circuit what the galvanometer shunt is to a galvanometer; by means of a slider the resistance of the deriving circuit, and consequently the magnitude of the current diverted into the principal circuit, can be increased or diminished at will. One of the chief uses to which the rheochord is put is in rheotome observations, to compensate any current of injury that may accidentally be present in muscle or nerve.

**Electrical resistance** is measured by means of a *Wheatstone bridge*, the principle of which is as follows. The current of a battery is sent through a branched circuit containing three known resistances, A, B, C, the unknown resistance,  $x$ , and a galvanometer placed in the bridging wire. Two of the resistances, A and B, are fixed; the third resistance, C, is variable, and used as the measuring resistance.

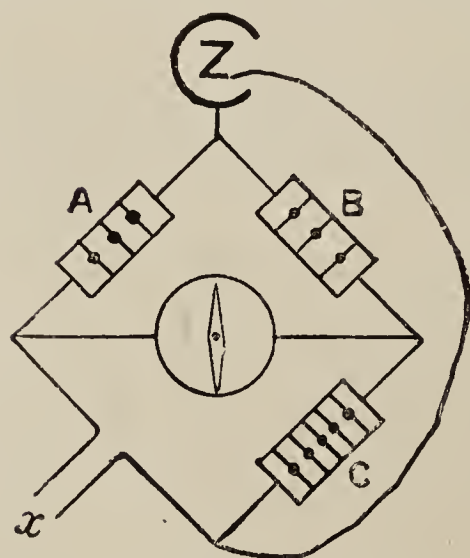


FIG. 123.—WHEATSTONE BRIDGE.

When the relation  $\frac{x}{C}$  is equal to

$\frac{A}{B}$  there is no current through the galvanometer, which therefore acts as a balance

indicating when C has been made too

large or too small. The equality  $\frac{x}{C} = \frac{A}{B}$

having been found by adjustment of C, the value of the unknown resistance,  $x$ , is found by simple calculation from the known values, A, B, and C; thus  $x = C \times \frac{A}{B}$ . If A and B are taken of equal resistance, the value of  $x$  is equal to that of C, as found by trial. These

facts should be realised by determinations of the resistance of muscle and of nerve as ordinarily arranged in physiological experiments. The following very approximate numbers will serve to give an idea of the resistances commonly dealt with in such experiments :—

Resistance of du Bois-Reymond's unpolarisable electrodes	. 10,000 ohms.
„ of d'Arsonval's silver electrodes . . . . .	. 100,000 „
„ of frog's gastrocnemius . . . . .	. 10,000 „
„ of 1 cm. frog's sciatic . . . . .	. 100,000 „
„ of human skin (according to moisture) . . . . .	. 1,000 to 100,000 „

The transverse resistance of nerve is about 5 times as great as its longitudinal resistance; that of muscle is about 10 times as great (Hermann). The difference is due to polarisation.

**Current diffusion—Distribution of potential.**—A current led through a bulky conductor, *e.g.* the human body, by two electrodes, *diffuses* throughout that conductor in accordance with Ohm's law. It is most concentrated—*i.e.* its density is greatest—along a line of least resistance between the two electrodes; it is less and less concentrated along curved lines further and further removed from this current axis. The nature of current diffusion and of the corresponding distribution of potential



FIG. 124.

will be best appreciated by the study of a simple case—a large vessel of salt solution into which the current of a Daniell cell is led by a fixed pair of silver chloride electrodes touching the surface, and from which



current can be led off by a movable pair of electrodes to a galvanometer. It will be found by trial that most current passes in the straight line  $+ -$ , less and less current along  $c_1, c_2, c_3$ , &c.; that if both leading-off electrodes are on points of the line  $OO$ , or on any two points of the lines  $+_1, +_1, +_1$ , or,  $+_2, +_2, +_2$ , &c., no current passes; that a greater current is obtained if the leading-off electrodes are on opposite sides of the line  $OO$  than if they are on the same side. Using the leading-off electrodes in connection with an electrometer instead of a galvanometer, we shall be able to test the electrical pressure differences between various points of the surface, and shall find that all that surface on the  $+$  side of  $OO$  is positive to all that surface on the  $-$  side of  $OO$ ; that the greatest difference is at and near the two points  $+$  and  $-$ ; that there is no difference between any two points on  $OO$ , or on any curved line,  $+_1, +_1, +_1$ ;  $-_1, -_1, -_1$ , &c. By these experiments we shall have defined the line  $+ -$  as the *current axis*;  $c_1, c_2, c_3$ , as *lines of current diffusion*;  $OO$ , cutting the middle of  $+ -$  at right angles, as the *equator*, or line of zero pressure, dividing an area of *positive* potential from an area of *negative* potential;  $+_1, +_1, +_1, -_1, -_1, -_1$ , &c., as *equipotential lines*, i.e. as lines all points of which are at the same electrical level. We shall find a very simple illustration of these elementary principles when we come to the study of the electrical variations of the human heart (p. 394).

**The galvanometer** (Thomson's).—The presence and the direction of an electrical current are ascertained by a *galvanoscope*, and its mag-

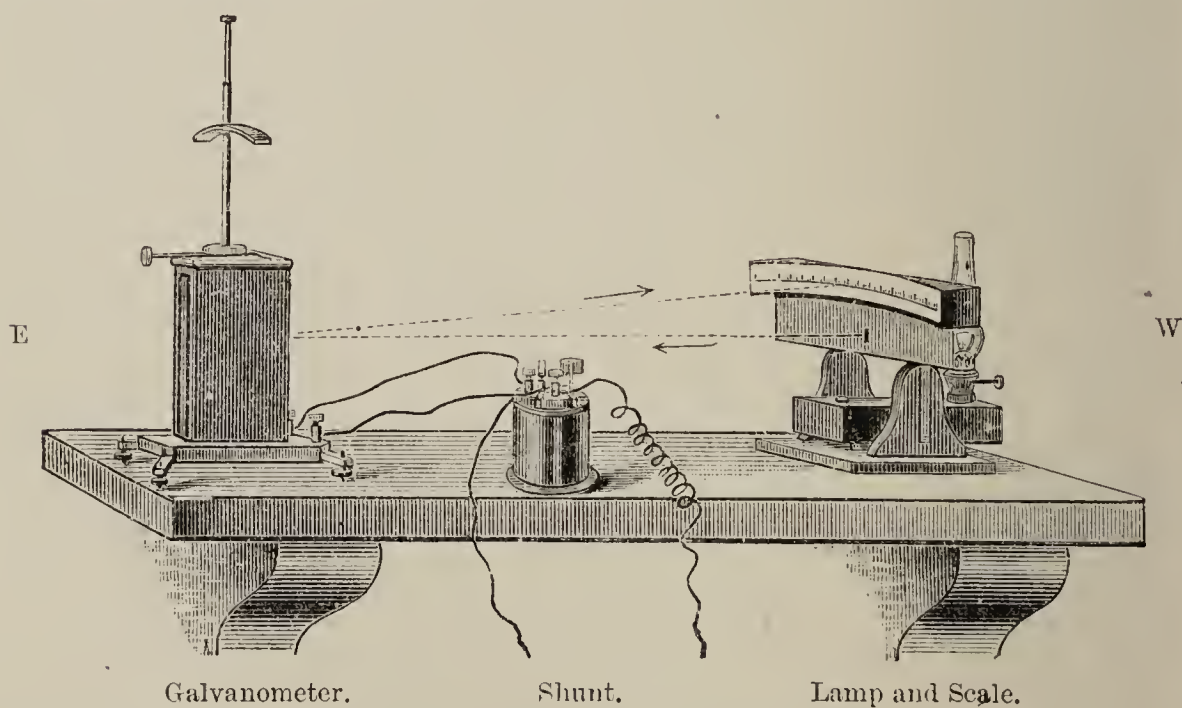


FIG. 125.—SIDE VIEW OF GALVANOMETER AND SHUNT, LAMP AND SCALE.

The galvanometer and scale are placed east and west, and appear as if viewed by an observer standing on the north side and looking south; the path of light is indicated by dotted lines. The essential parts concealed by the galvanometer case are diagrammatically given in fig. 126.

nitude by a *galvanometer*; a graduated *galvanoscope* is a *galvanometer*, and obviously any *galvanometer* may be used merely as a *galvanoscope*.

For clinical purposes it is sufficient to use a small instrument graduated from one to twenty milliamperes ; for laboratory purposes, *e.g.* for nerve-currents, an instrument at least 10,000,000 times more sensitive is required, owing to the magnitude of the resistances, and the smallness of the differences of potential to be dealt with. A galvanometer of this kind is of *high resistance* (5,000 to 20,000 ohms). When necessary its sensitiveness is reduced by the use of a *shunt*, which carries off  $\frac{9}{10}$  or  $\frac{99}{100}$  or  $\frac{999}{1000}$  of a current through a deriving circuit,  $\frac{1}{9}$  or  $\frac{1}{99}$  or  $\frac{1}{999}$  of the resistance of the galvanometer, and therefore leaving to pass through the galvanometer  $\frac{1}{10}$  or  $\frac{1}{100}$  or  $\frac{1}{1000}$  of the original current. For *thermo-electric* measurements, in which small differences of potential through a very small resistance are dealt with, a galvanometer of *low resistance* (less than 1 ohm) is employed. The principle upon which both these galvanometers depend is, that a suspended magnet surrounded by coils of wire is deflected when a current passes through the wire, the direction of the deflection and its amount indicating the direction and amount of the current. The suspended magnet (or system of magnets) is made very nearly ‘astatic,’ *i.e.* not to set too strongly towards the magnetic pole, and can be made more or less astatic, and therefore more or less sensitive, by means of a separate magnet, which can be adjusted to neutralise the ‘set’ of the suspended magnet more or less completely, or if necessary be reversed so as to increase it. The movements of the suspended magnet are shown greatly magnified by means of a light mirror which reflects a beam of light on to a horizontal scale. If desired, the varying positions of the spot can be photographed upon a travelling sensitive surface placed behind a slit in the scale (*e.g.* figs. 152, 163, 177, 182, 199, 201).

In Wiedemann’s galvanometer the adjustments are effected by movable coils, fine wire coils being used for currents through high resistance, coarse wire coils for currents through low resistance ; the coils can be set nearer or further from the suspended magnet, the deflections of which are read through a telescope.

In d’Arsonval’s galvanometer the coil is suspended between two strong magnets, and is itself deflected by the currents under observation.

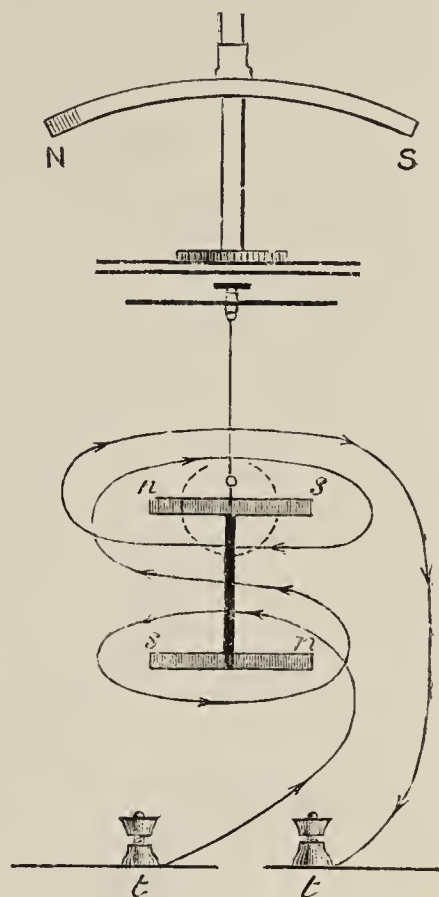


FIG. 126.

Astatic couple of magnets *n s, s n*, suspended by a silk fibre and carrying a mirror (indicated by the dotted circle); the surrounding line and arrows indicate the disposition of the coils ; *n s* is the neutralising or controlling magnet. All these parts are represented as if viewed by an observer standing west, *i.e.* in the position of the lamp in fig. 125.



The deflections are observed by a telescope, or (with sufficiently strong currents from one millième upwards) can be recorded by means of a light pen fixed to the coil on a smoked cylinder moving round a horizontal axis. Such records are particularly desirable in experiments on the excitability of human muscle and nerve.

The suspended magnet of an ordinary galvanometer, after it has been set in movement, comes to rest by a series of diminishing oscillations above and below its position of rest. In such case the instrument is said to be '*undamped*' or '*partially damped*,' and an instrument in which the unavoidable damping due to the resistance of the air is made as small as possible is called a *ballistic* galvanometer. If, on the contrary, the damping is increased, the magnet is rendered '*dead-beat*' or '*aperiodic*,' i.e. its oscillations are suppressed so that it does not swing beyond its steady deflection by a current, nor beyond its zero with cessation of current. In such case the movement by which the magnet takes up its position occurs more slowly than in that of an undamped magnet, and with a gradually increasing slowness; the time occupied by this movement is called the '*falling time*.' With an undamped magnet the time occupied by one oscillation to and fro is its '*period*,' and the relation between the magnitudes of two successive oscillations is the '*decrement*.'

The galvanometer is an indicator of current, but by adopting the method of **compensation** it becomes an indicator of potential, being in

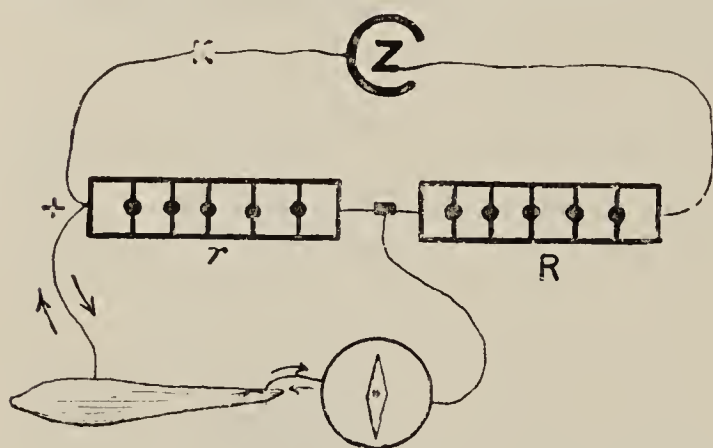


FIG. 127.—MEASUREMENT OF POTENTIAL BY COMPENSATION.

this case used as a balance to show the equality of two opposite potentials—the galvanometer circuit being then free of current. The simplest way of measuring the potential difference<sup>1</sup> at any two points of muscle or nerve, &c., is as follows (fig. 127): A Daniell cell is connected with the two ends of a rheostat, divided into two parts, one of

lower resistance,  $r$ , the other of higher resistance,  $R$ . With the high external resistance, the P.D. at the two terminals is equal to the full E.M.F. of the cell (i.e. 1 Dan. or about 1.1 volt), or differs from it by a negligible fraction (i.e.  $\frac{\text{internal resistance}}{\text{external resistance}}$ ), and the P.D. at any two points of the circuit varies directly as the resistance between these points. The P.D. at two points of a muscle is ascertained by finding the resistance,  $r$ , at which it is balanced (i.e. no current through the galvanometer); it is then equal to  $r : r + R$ . For example, if the

<sup>1</sup> 'Potential difference,' 'electromotive force,' are shortly expressed by the initial letters P.D., E.M.F.

balance is obtained with  $r=250$  ohms, and  $r+R=7,500$  ohms, the muscle potential is  $250 : 7,500$  or  $0.33$  Dan. The principle is precisely the same as that of the rheochord, but the total resistance in the battery circuit is much greater.

An indispensable condition to be observed whenever the electrical excitability of exposed nerve or muscle is to be exactly studied, and still more whenever currents from living tissues are to be led off to a galvanometer, is the employment of **unpolarisable electrodes** (du Bois-Reymond). Their construction is as follows:—A carefully-amalgamated zinc rod dips into a saturated solution of zinc sulphate, which in turn communicates with a plug of china clay made up into a paste with

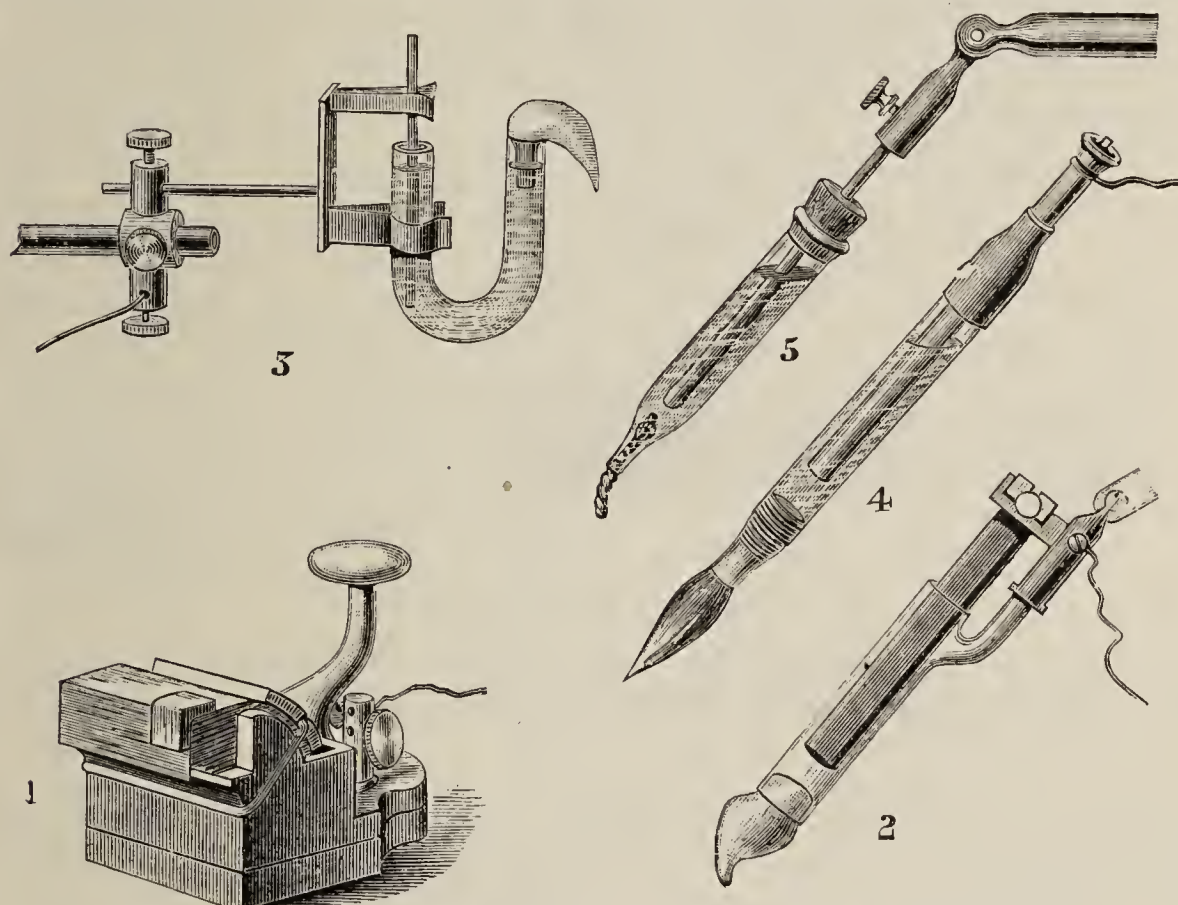


FIG. 128.—SEVERAL MODELS OF UNPOLARISABLE ELECTRODES.

1 and 2, du Bois-Reymond's; 3, Burdon-Sanderson's; 4, von Fleischl's; 5, d'Arsonval's.

In 1, 2, 3, and 4 the component parts are zinc, zinc-sulphate, and saline clay; in 5 a silver rod coated with fused silver chloride dipping in normal saline contained in the tube from which a thread projects.

normal saline; a glass tube shaped according to requirements, and fixed in a suitable holder, contains the several constituents of such an electrode, which has the following qualities: it is unpolarisable by weak currents; it is not itself a source of electromotive force; it is of high resistance; it can be applied to living tissues without appreciably injuring them. A pair of unpolarisable electrodes should be tested before use by bringing their plugs into contact while they are connected with the galvanometer; they should then give little or no current. A more convenient (though less perfect) unpolarisable electrode is that of d'Arsonval; it consists of a silver rod coated with fused silver chloride dipping into a tube filled with normal saline.



The **rheotome** (Bernstein, Hermann) is an instrument by which a series of stimuli can be led into muscle or nerve, and the consequent series of excitatory effects led off to a galvanometer, during definite periods at regular intervals after stimulation. This is in principle

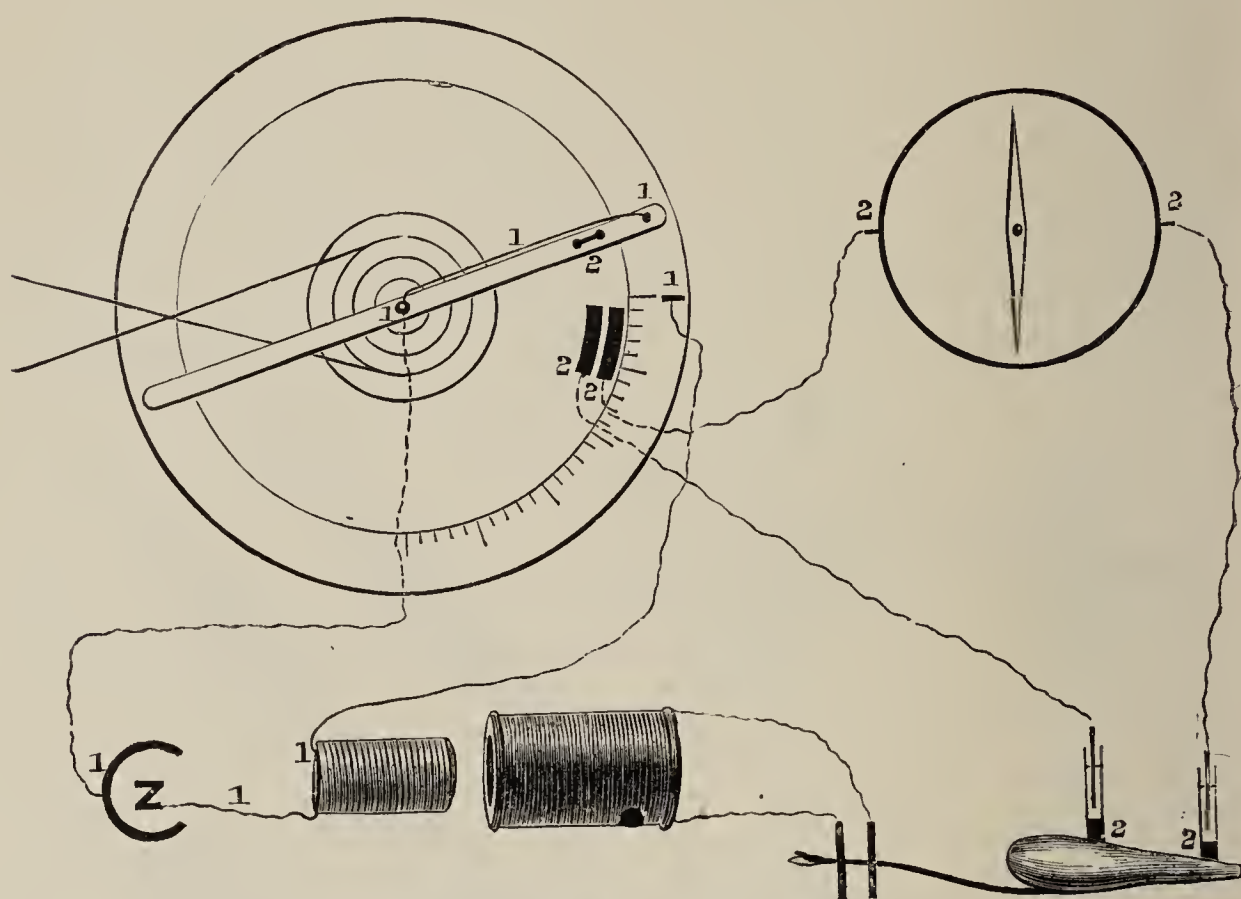


FIG. 129.—SCHEMA OF A RHEOTOME.

1 1 1 1 is the primary or excitation circuit; 2 2 2 2 is the galvanometer circuit. Any current from the quiescent muscle would be balanced by a compensator, as shown in fig. 122.

effected by a revolving bar carrying two contacts, one in the primary or exciting circuit, 1 1 1, one in the galvanometer or electrode circuit, 2 2 2 (the latter being in the form of a metal horseshoe ending by two brushes of fine wires which rub against the metal plates connected with the galvanometer and leading-off electrodes). If the bar revolves once a second, and completes the galvanometer circuit by a pair of metal pieces occupying  $\frac{1}{100}$  part of the circle, the circuit is closed at each second for  $\frac{1}{100}$  sec. The circle carrying the primary contact is movable round the circle carrying the galvanometer contact; it can be set so that the two contacts are made simultaneously, or so that the galvanometer contact, lasting  $\frac{1}{100}$  sec., is made  $\frac{1}{100}$ ,  $\frac{2}{100}$ ,  $\frac{3}{100}$ , &c. sec. later than the primary contact. We can thus obtain a repetition of effects, and 'tap' the excited muscle or nerve at each revolution—*i.e.* each second—for a period of  $\frac{1}{100}$  sec. immediately after each stimulus, or  $\frac{1}{100}$ ,  $\frac{2}{100}$ ,  $\frac{3}{100}$  sec. later. Combinations suitable for various purposes are obtained by varying the rate of revolution, the length of the galvanometer contact and its angular distance from the exciting contact; *e.g.* if the bar revolves 10 times a second and makes contact for  $\frac{1}{20}$  part of the circle, the circuit is closed for  $\frac{1}{200}$  sec.; if for  $\frac{1}{50}$  part of the circle, the circuit

is closed for  $\frac{1}{500}$  sec. Any injury or other current from the tissue must be kept exactly compensated during experiment.

**Thermo-electric measurements.**—A thermo-electric couple is the surface of junction of two different metals that form a completed circuit. If the junction is heated, an electrical current passes across the junction and around the circuit, as is seen by the deflection of a galvanometer placed in the circuit. Different pairs of metals act more or less strongly when equally heated. The couples usually employed are bismuth and antimony, or german silver and iron; the arrows indicating the direction of the current through the heated junction. Within certain limits, strength of current is in direct proportion with temperature of junction; it is thus possible to read temperature by means of the galvanometer (which indicates strength of current).

If two similar couples are in circuit with a galvanometer (as shown in the second diagram), and both junctions are

heated equally, no current passes through the galvanometer, the current through one couple from B to A being balanced by the opposed current through the other couple from B to A. But if there be any inequality of temperature between the two couples, then a current passes, the direction of the deflection indicating which is the warmer couple. We may thus readily determine differences of temperature by plunging needle-shaped thermo-electric couples into the tissues. Or, having plunged two needles symmetrically into a frog's gastrocnemius of each limb, we may excite first one, then the other, sciatic nerve, and observe deflections first in one, then in the opposite, direction, indicating the production of heat in the muscular contraction, first on

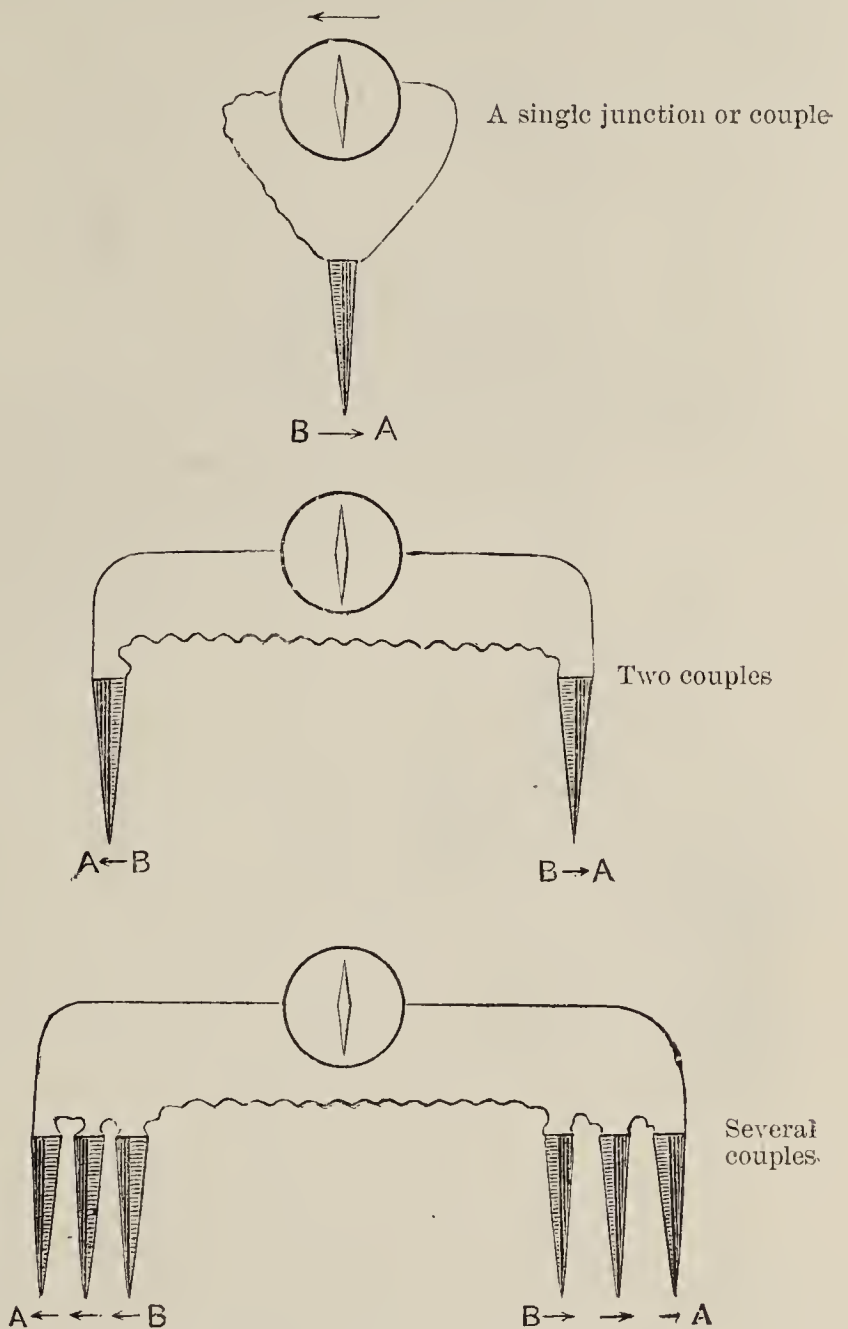


FIG. 130.—SCHEMA OF THERMO-ELECTRIC COUPLES.



one, then on the opposite, side (fig. 152). For very delicate observations two series of couples are used as thermopiles.

**The capillary electrometer (Lippmann).**—A glass tube drawn out at one end to a fine bore (20 to 30  $\mu$ ) is filled with mercury and connected with an apparatus by means of which the pressure can be raised or lowered. The capillary end of the tube dips into 10 per cent. sul-

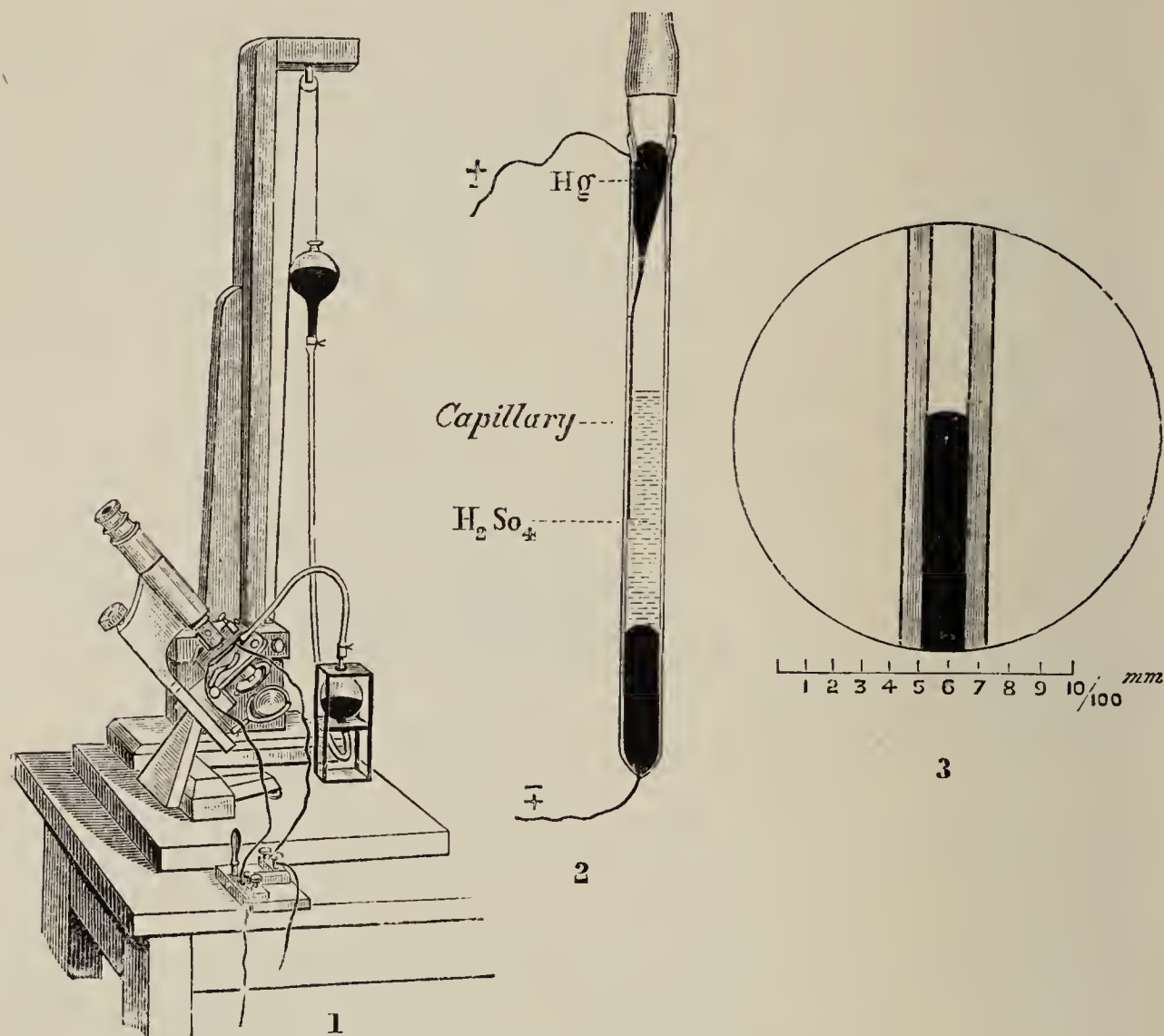


FIG. 131.—LIPPMANN'S CAPILLARY ELECTROMETER.

1. Pressure apparatus and microscope, on the stand of which the capillary tube is fixed.

2. Capillary tube dipping into  $\text{H}_2\text{SO}_4$  in a surrounding tube, and in connection with pressure apparatus (the mercury in the lower part of the surrounding tube serves only to establish connection with the platinum wire).

3. The capillary tube and column of mercury as seen in the field of the microscope. (Scale in  $\frac{1}{100}$ ths mm.)

phuric acid. Two platinum wires fused through the glass establish connection with the mercury and with the sulphuric acid respectively. By means of the pressure-apparatus mercury is forced into the capillary and adjusted until it takes up a position of rest in the field of a microscope. The pressure of the mercury and the capillarity of the tube are in equilibrium; the lower surface of the mercury is in a state of tension, which is very easily increased or diminished by variations of electrical potential. The instrument is, in fact, an exceedingly delicate electrical manometer; a rise of electrical pressure on the mercury

side, or a fall of electrical pressure on the sulphuric acid side, causes the mercury to move towards the point of the capillary; a fall of electrical pressure on the mercury side, or a rise on the sulphuric acid side, causes the mercury to recede from the point of the capillary. The instrument accordingly is an indicator of 'potential' or 'pressure,' not of 'current.' Its delicacy is such that it will react to as little as  $\frac{1}{40000}$  volt. It offers, moreover, the following advantages:—the indications are practically instantaneous, free of lost time, and of after-oscillation; the resistance in circuit is immaterial; unpolarisable electrodes may for most purposes be dispensed with.

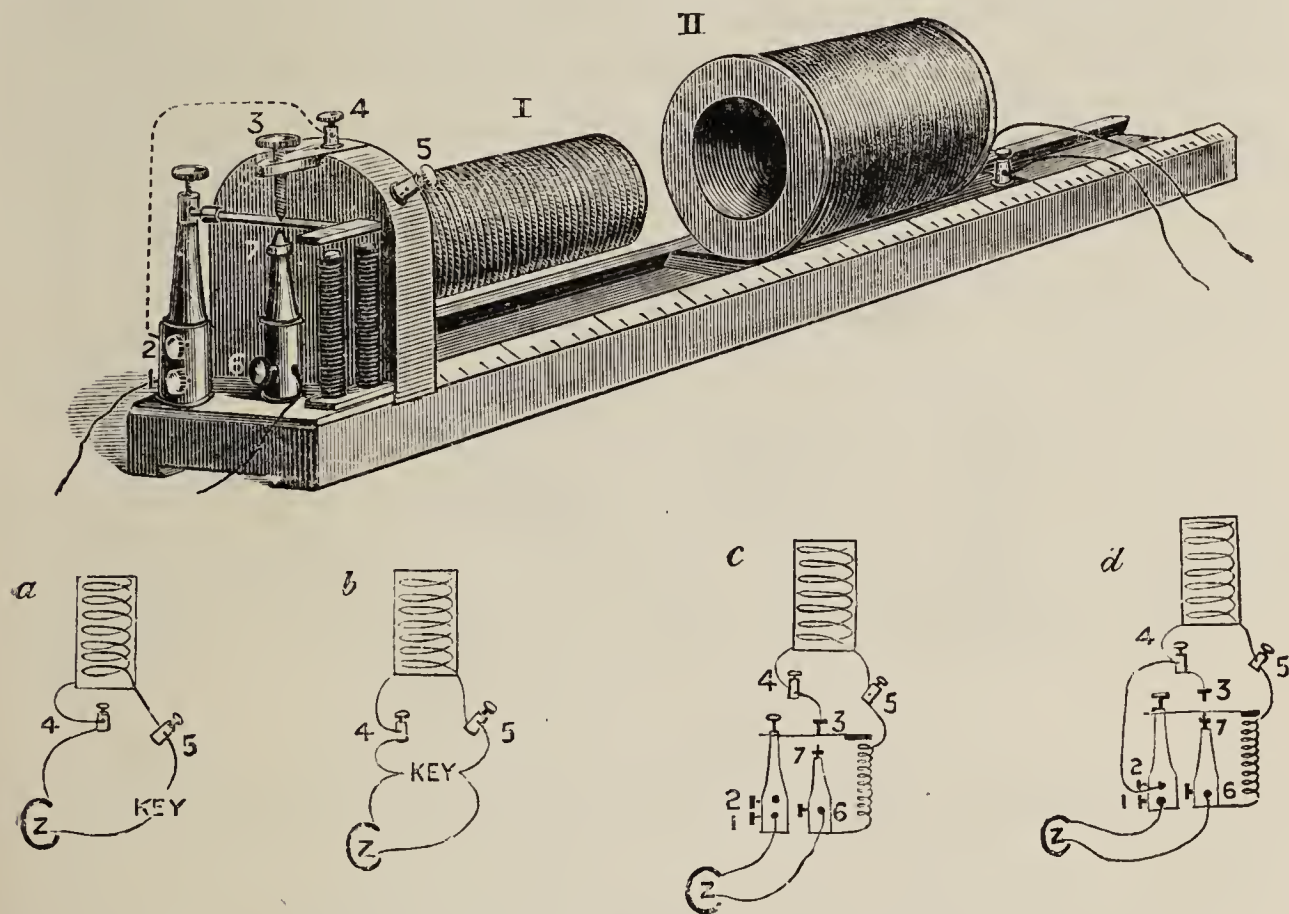


FIG. 132.—DU BOIS-REYMOND'S INDUCTION APPARATUS.

The numbers 1 to 7 indicate the terminals and contact screws connected with the primary coil, I.

For single shocks the two battery wires are to be connected with the terminals 4 and 5, which are at the two ends of the primary wire.

(a) *Unmodified shocks* are obtained when a key is used to interrupt one of the wires.

(b) *Reduced shocks* are obtained when a key is used for short-circuiting the primary wire.

(c) For repeated shocks (ordinary) the two battery wires are to be inserted at 1 and 6. The circuit now includes the spring interrupter and the wire of the electro-magnet by which the circuit is made and broken at the contact screw 3; the contact screw 7 is kept out of use by being lowered.

(d) For repeated shocks (modified) the battery wires are left, as before, at 1 and 6. A short, thick sidewire is placed between 2 and 4. The contact screw 3 is raised out of range of the spring, and the contact screw 7 is raised until it comes within range of the spring.

The electrode wires are in each case connected with two terminals (not seen in figure) forming the two ends of the secondary coil, II.

The induction apparatus of du Bois-Reymond is the electrical instrument most often used in laboratories and in hospitals and the



main principle upon which its working depends is that a current on commencing to flow, and on ceasing to flow through one coil of wire, induces an opposite current of short duration and great tension in an adjacent but otherwise independent coil. The apparatus consists of (1) a cell or battery that supplies electromotive force, (2) a primary or thick wire coil, (3) a secondary or thin wire coil. The cell and connected primary coil form the primary circuit, which contains in addition a key or interrupter; the secondary coil, connected with the nerve or muscle by wires and electrodes, forms the secondary circuit.

When, by closing the key, the battery current is made through the primary coil, a make induction current is induced in the secondary coil. During the passage of the primary current nothing happens in the secondary coil. When, by opening the key, the battery current through the primary coil is broken, a break current is induced in the

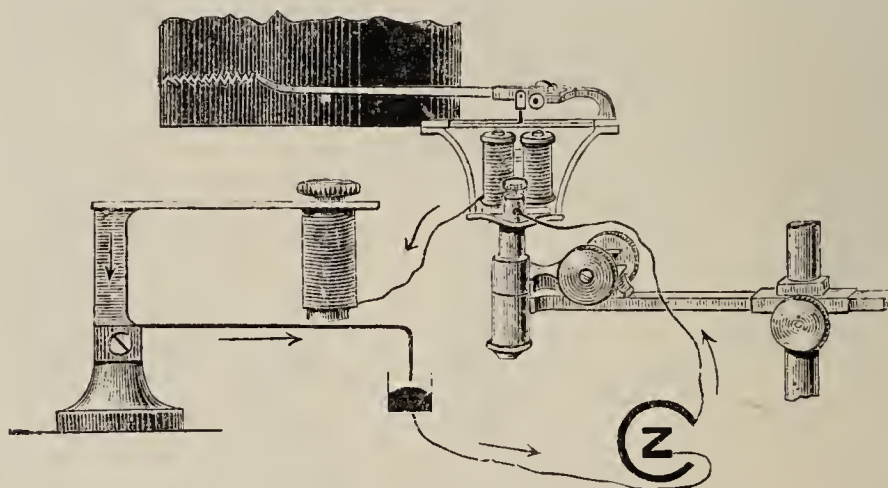


FIG. 133.—CHRONOGRAPH, OR TIME-MARKER.

Composed of battery, vibrating reed (Page's), signal (Pfeil's).

secondary coil. The make current and the break current in the secondary coil are of opposite directions, and—as should be appreciated by placing the electrodes on the tongue—of unequal intensities. The break shock is sharper than the make shock; a difference that should serve to remind us not to confuse the make and break currents induced in a secondary coil with the make and break of a battery current directly led through a nerve or muscle (p. 346). The disparity (as fully explained in the Practical Guide) depends upon extra currents due to self-induction, and can be remedied by the Helmholtz side-wire (fig. 132). The strength of currents induced in the secondary coil by the make and break of the inducing current in the primary coil, depends upon the distance between the two coils; it increases as the coils are brought closer together, diminishes as they are removed further apart; thus by sliding the secondary towards or from the fixed primary coil the strength of this form of excitation can be conveniently varied. With a simple make and break key in the primary circuit, single shocks are obtained. Rapidly repeated series of shocks—*i.e.* Faradisation or tetanising currents—are obtained by the interpola-

tion of an automatic interrupter or vibrating key, which when in action is the cause of the noise usually associated with induction coils.

**Electrical chronographs.**—Measurements of time are in constant requisition for physiological purposes, and among the various means adopted an electrical method is the commonest, as we can thus most easily take a chronographic record simultaneously with the movement recorded. The component parts of electrical time-markers are a small electro-magnet with a recording lever attached to the armature, and with its coil in the circuit of a cell and interrupter. The interrupter may be a tuning-fork or a reed of given vibration-frequency, with a platinum point, which by means of a second electro-magnet is kept vibrating to and from a clean surface of mercury, making and breaking circuit, attracting and releasing the chronograph armature, and thus marking time in corresponding fractions of a second. The time-markers in ordinary use are those of Despretz and of Pfeil. If the apparatus is used as a signal, the electro-magnetic interrupter is replaced by a key, and the lost 'time' of the signal itself should be determined; this is usually from  $\cdot 0005$  to  $\cdot 0010$  second.

A spring, held in a clamp so that by adjusting its length its vibration-frequency can be altered at will, forms a simple and convenient interrupter when placed in the primary circuit of a coil so as to vibrate in and out of a mercury cup in the same circuit; such an interrupter can easily be graduated to give from ten to fifty stimuli per second.

## MUSCLE

The muscles of animals, anatomically as well as physiologically considered, are divisible into two separate and distinct systems—(1) those which form the chief bulk of the walls of the hollow viscera and of the blood-vessels; (2) those which are attached to the bones. Muscles of the first kind are not subject to the will, and are therefore characterised as *involuntary*; they are composed of long nucleated cells which are not striped; hence such muscle is characterised as *smooth* or *non-striated*. Muscles of the second kind are under the control of the will, and are therefore called *voluntary*; they are composed of fibres which are transversely striped; hence such muscle is also characterised as *striated*. Between these two typical kinds—the smooth involuntary and the striated voluntary—a third kind must be reckoned, viz. cardiac muscle, which is intermediate in its anatomical and physiological characters. Voluntary striated muscle is further distinguishable in many animals into two varieties, viz. *pale* and *red*. The pale variety is the more highly organised; it exhibits more marked striation and less abundant nuclei than the red variety.



Involuntary muscle is chiefly supplied by non-medullated nerve-fibres, but also receives a smaller proportion of medullated fibres. The actual termination of nerve-fibre in involuntary muscle is not known with certainty; abundant networks of fibres are to be seen, and in connection with them ganglionic cells. Voluntary muscle is chiefly supplied by medullated nerve-fibres, which form definite structures at their junction with the muscular fibres, the most characteristic of these being motor end-plates.

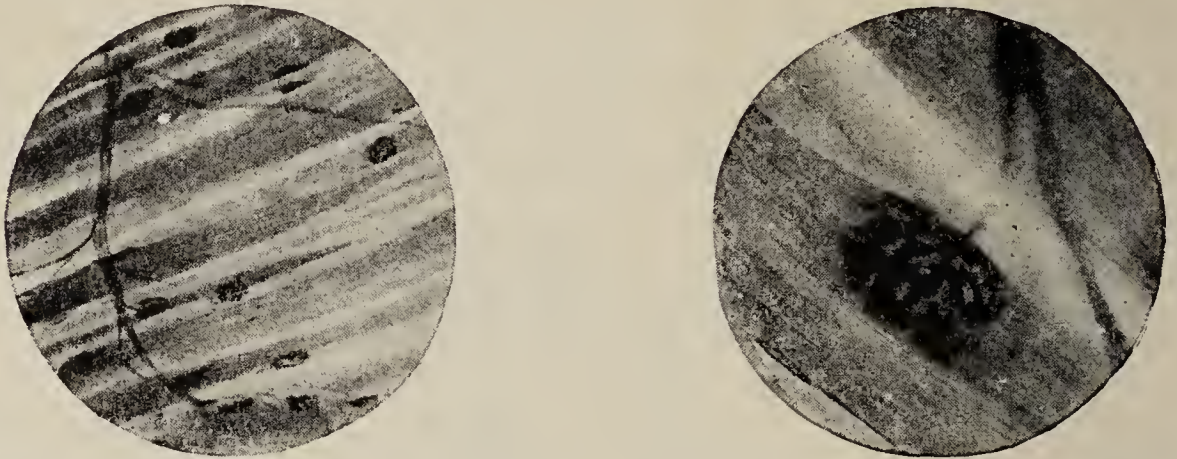


FIG. 134.—MOTOR NERVE-ENDINGS IN STRIATED MUSCLE, UNDER LOW AND HIGH POWERS.  
(Photographed from a gold preparation by Mr. F. S. Lloyd.)

**Histological characters of resting, of contracting, and of rigid muscle.**—The ultimate structure of a striped muscle-fibre has been for long a vexed question. Upon one important point all observers are agreed, viz. that the sarcolemma, or sheath of the fibre, contains contractile solid particles (sarccous elements) and a fluid non-contractile portion. That the fibre is not transversely divided by a series of partitions (Krause's membrane) is considered to be proved by Kühne's observation of a nematode worm moving freely in a living muscle-fibre without meeting with any apparent assistance until coagulation had occurred. By microscopic observation of a tetanised muscle-fibre, or of a muscle-fibre fixed in a state of partial contraction by osmic acid, it has been found that the shortening and thickening of the fibre are attended with an obvious approximation of its transverse striæ, and with a by no means obvious reversal of the optical relations of the striæ, the dark broad band of the resting muscle becoming the light narrow band of the contracting muscle. Viewed by polarised light, there is no such transposition; the dark band of an uncontracted fibre and the light band of a contracted fibre are both doubly refracting (anisotropous) and appear as light bands in a dark field. We must, however, make the express reservation that these observations refer in both cases to dead muscle, and

that it is nowise certain that the partially contracted parts of muscles fixed by osmic acid give the true picture of a natural contraction. And, as a matter of fact, it is only dead muscle

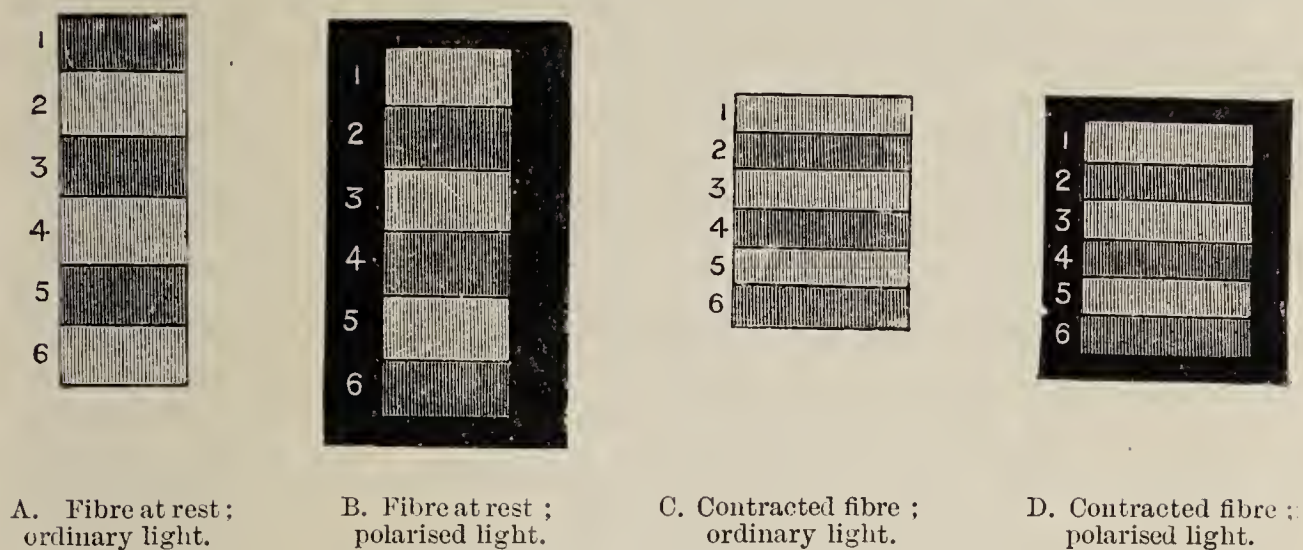


FIG. 135.

The accompanying very diagrammatic figure is intended only as a guide to the description given in the text, and not in any degree to imitate the actual appearances. The odd numbers 1, 3, 5, point to contractile substance, dark in A, light in C; light in B and D (*i.e.* doubly refracting or anisotropic). The even numbers, 2, 4, 6, point to non-contractile fluid, light in A, dark in C; dark in B and D (*i.e.* singly refracting, or isotropous).

which exhibits under polarised light the regular and well-marked alternation of doubly-refracting light band and singly-refracting dark band; in living muscle almost the whole of the substance is doubly refracting; singly refractile substance in the form of lines and dots is relatively scanty.

**Chemical composition of muscle; its alteration by activity and in rigor.**—Muscle—or, to call it by its popular names, flesh or meat—is mainly composed of water and of proteid; the elementary composition of dry flesh is thus closely similar to that of dry proteid or of dry blood (*vide* p. 10). Besides its proteid constituents, muscle contains a comparatively small quantity of other nitrogenous bodies, of inorganic salts, and of carbohydrates; a variable amount of fat is usually associated with it, but is not, properly speaking, a constituent of pure muscle. The average composition of muscle is as follows:—

Water	.	.	.	.	.	.	.	.	.	.	.	77.5
Proteid	.	.	.	.	.	.	.	.	.	.	.	20
Other nitrogenous bodies (creatin, xanthin)	.	.	.	.	.	.	.	.	.	.	.	1
Salts (phosph. and K)	.	.	.	.	.	.	.	.	.	.	.	1
Carbohydrates (glycogen, inosit, lactic acid, sugar)	.	.	.	.	.	.	.	.	.	.	.	.5

The red colour of muscle is partly due to blood-pigment, partly to a special pigment—myohæmatin (MacMunn); ferments are also present—viz. the myosin ferment and a peptic ferment.



We have to study (1) the chemical differences between living and dead muscle; (2) the chemical differences in dead muscle, which before death was at rest or in forced activity.

The consistence of a living irritable muscle differs from that of dead muscle which has entered into 'rigor mortis'—the former is soft and elastic, the latter is firm and very imperfectly elastic—'doughy.' The difference depends upon the fact that living muscle consists of separate solid and liquid parts—the sarcous elements and muscle-plasma—while in dead muscle the liquid part (plasma) has undergone coagulation and has yielded myosin. The rigor or coagulation of muscle is, indeed, closely analogous with the coagulation of blood: it depends upon the conversion of a body present in muscle-plasma (myosinogen) into myosin, just as the coagulation of blood depends upon the conversion of a body present in blood-plasma (fibrinogen) into fibrin, and it is probable that in the case of muscle, as in that of blood, the conversion is effected by the agency of a ferment. The myosin coagulum exhibits the following remarkable difference from the fibrin coagulum: both are soluble in 10 per cent. solution of sodium chloride, the first-named body the more readily; the dissolved myosin yields a renewed coagulum when its salt solution is diluted: not so the dissolved fibrin (Halliburton). The myosin coagulum having formed is thus easily unmade and remade out of the body; *perhaps* this is also the case in the body.

*Preparation of muscle-plasma.*—The coagulation of muscle-plasma is delayed by cold. The fresh, irritable muscles, preferably of frogs, are accordingly frozen, minced, pounded, and pressed, all instruments used being cooled; a syrupy alkaline liquid is expressed, which is coagulable at ordinary temperatures or on dilution with cold distilled water. This liquid is *muscle-plasma*, and the body that separates from it when coagulation occurs is *myosin*. The fluid that remains after removal of the coagulum is *muscle-serum*, which is acid, and contains in solution (1) muscle-globulin, coagulable at a temperature of 63°; (2) muscle-albumin, coagulable at 73°.

*Preparation of myosin.*—If muscle-plasma is allowed to fall drop by drop into distilled water, myosin coagulates at once, and is obtained in the form of little balls. Myosin may also be obtained from rigid muscle in which it has already formed, by pounding with sodium chloride, then adding water to make a 10 per cent. solution of the amount of salt used; myosin is soluble

in this, and is precipitated by pouring the solution thus obtained into a considerable quantity of water, or by dialysis. Myosin belongs to the class of globulins, and is as such insoluble in distilled water, soluble in solutions of neutral salts ( $\text{NaCl}$ ,  $\text{MgSO}_4$ ,  $\text{Na}_2\text{SO}_4$ ); in solution it coagulates when heated to between  $55^\circ$  and  $60^\circ$ .

*Sarcolactic acid*.—Living resting muscle has a neutral or alkaline reaction; dead rigid muscle has an acid reaction; muscle-plasma is alkaline, muscle-serum is acid. The difference is due to *sarcolactic acid* produced during rigor, *i.e.* accompanying though not necessarily caused by the conversion of myosinogen into myosin. We have already seen that the rigor of muscle is accompanied with a production of carbon dioxide (pp. 136–7); we are about to learn that the same two changes also accompany the normal activity of muscle. But whereas in rigor the acidification is the more prominent phenomenon, in normal action the chief event is the production of  $\text{CO}_2$ ; these considerations suggest as probable that the lactic acid ( $\text{C}_3\text{H}_6\text{O}_3$ ) produced in muscular katabolism is a stage towards the production of  $\text{CO}_2$ ; in full action muscle may produce lactic acid, and then decompose it further into  $\text{CO}_2$ ; in declining action muscle may produce lactic acid, and fail to complete the process.

*Inosit*, or muscle-sugar ( $\text{C}_6\text{H}_{12}\text{O}_6, 2\text{H}_2\text{O}$ ), is a minute and variable constituent of muscle. Unlike grape-sugar, it has neither rotating action on polarised light, nor reducing action upon Fehling's solution; it is a crystalline body that can undergo lactic but not alcoholic fermentation, and is now recognised, not to be a true carbohydrate, but to belong to a benzoic series (see p. 596).

*Rigor mortis* as manifested by the human body is of medico-legal interest, as yielding some indication of the time at which death may be presumed to have occurred. From observations on tetanised muscles and on hunted animals we learn that previous muscular activity hastens the advent of rigor; from observations on man—deaths by accident, or in hospitals, or in battle—we learn that the 'death-stiffening' begins earliest in a person surprised in active exertion at the time of death; on worn-out bedridden patients it begins early, is ill-developed, and soon passes off. It usually begins two or three hours after death, and lasts for two or three days, but soldiers have been found in attitudes indicative of its immediate onset, and it sometimes



does not commence before several hours after death. The order of its manifestation on the human subject is from above downwards, in the jaw, neck, arms, and legs. After a period varying from a few hours to a few days, rigor passes off, and in a reverse order, by resolution of the coagulated myosin, and now putrefaction sets in.

*Differences between active and resting muscle.*—There are only two known chemical differences between living resting and living active muscle, viz. the reaction and the amount of  $\text{CO}_2$ —living resting muscle is neutral or faintly alkaline in reaction, living active muscle is slightly acid, the acidity, like that of rigor, being due to sarcolactic acid. In its chemical effects, activity is thus comparable with partial death, and we may add to the comparison by the observation that whereas death is attended by the complete and final coagulation of myosin, prolonged activity is marked by an incompleteness of relaxation, which *may be*—though it cannot be *proved* to be—due to a temporary coagulation of myosin. The similarity of the electrical change, which accompanies the excited and the dying states, and the evolution of heat which accompanies contraction and rigor, may be alluded to as further points of resemblance. Fick offers as the most probable hypothesis to account for the phases of contraction and relaxation (1) a production of lactic acid and of myosin during the former, (2) a production of  $\text{CO}_2$  and a resolution of myosin during the latter. In pursuing our examination beyond this stage, it is to be remembered that the chemical differences between active and inactive muscle relate in each case to muscle that is dead, having necessarily been killed in the process of analysis. In correspondence with the statement just made, that muscular activity is attended with an evolution of  $\text{CO}_2$ , it is found that the  $\text{CO}_2$  collected by means of the gas-pump (*a*) from rigid previously active muscle, (*b*) from rigid previously resting muscle, is of greater amount in the second case than in the first; *i.e.* previously active muscle contains less  $\text{CO}_2$ -yielding substance than previously resting muscle. Further than this only some isolated statements have been made by various observers, to the following effect:—

1. Previously active muscle yields a smaller amount of water-extractives but a larger amount of alcohol-extractives than previously resting muscle (Helmholtz).

2. Pyrogallie acid is more readily oxidised by an aqueous extract of resting muscle than of tetanised muscle. Sulphate of

indigo is more readily reduced by an aqueous extract of tetanised muscle than of resting muscle (Grützner).

3. Resting muscle contains more glycogen; tetanised muscle contains more sugar (Nasse).

**The properties of muscle.**—The physiology of muscle is almost entirely derived from the study of voluntary muscle, and the considerations upon which we are now entering relate almost exclusively to voluntary muscle.

Living muscle possesses two chief properties: (1) It is extensible and elastic; (2) it is excitable and contractile.

*Extensibility and elasticity; viscosity.*—Muscle can be stretched beyond its normal length, and can recover its original length when the extending force is removed, unless the extension has been excessive. Extensibility is demonstrated by the elon-

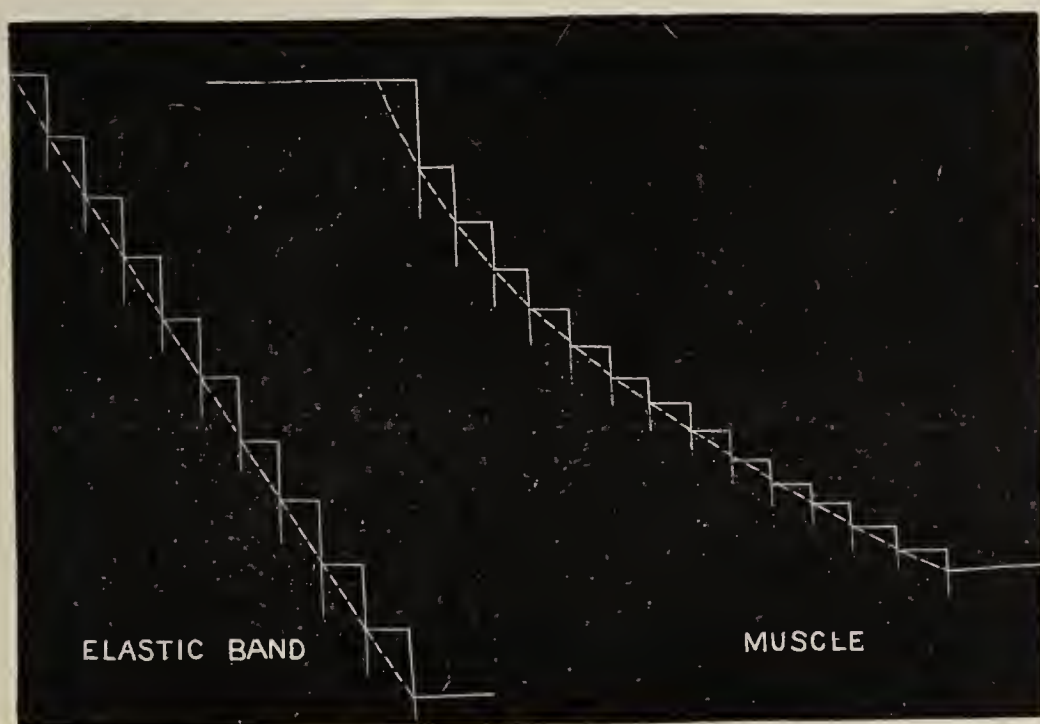


FIG. 136.

gation of a muscle fixed at one end and weighted at the other; elasticity is demonstrated by the shortening when the weight is removed.<sup>1</sup> The magnitude of these alterations is conveniently studied by means of a long light lever, which is attached to the lower end of the muscle, and records the changes of length upon

<sup>1</sup> Considerable confusion still prevails in the use of the term 'elasticity.' Sometimes it is improperly used to denote the 'yieldingness' of a body, which, after it has been stretched beyond its length, returns to its original length. This use of the term is incorrect; extensibility is the correct word to use. Sometimes it is used to denote resistance to stretching; but this is inconvenient, for in this sense of the term an inextensible body has great elasticity. It is preferable to limit the term to express the force with which a stretched or compressed body tends to return to its original dimension.



a smoked surface. A frog's gastrocnemius so disposed, and extended successively by weights of 10, 20, 30, 40, &c. grammes, will show for each successive equal increment of 10 grammes a diminishing increase of length; *i.e.* the elongation increases with the increased weight, but in a diminishing ratio; for instance, the elongation when the weight is increased from 20 to 30 is less than when the weight was increased from 10 to 20; it is less still when the weight is increased from 30 to 40, and so on. In this respect the extensibility of muscle resembles that of arteries, and differs from that of substances such as caoutchouc or indiarubber, which give a series of elongations nearly proportional to the weights used. The recovery of length dependent on elasticity is observed in either case by the successive removal of the weights. It will be noticed in the case of muscle that each recovery of length as each weight is removed is greater than the preceding

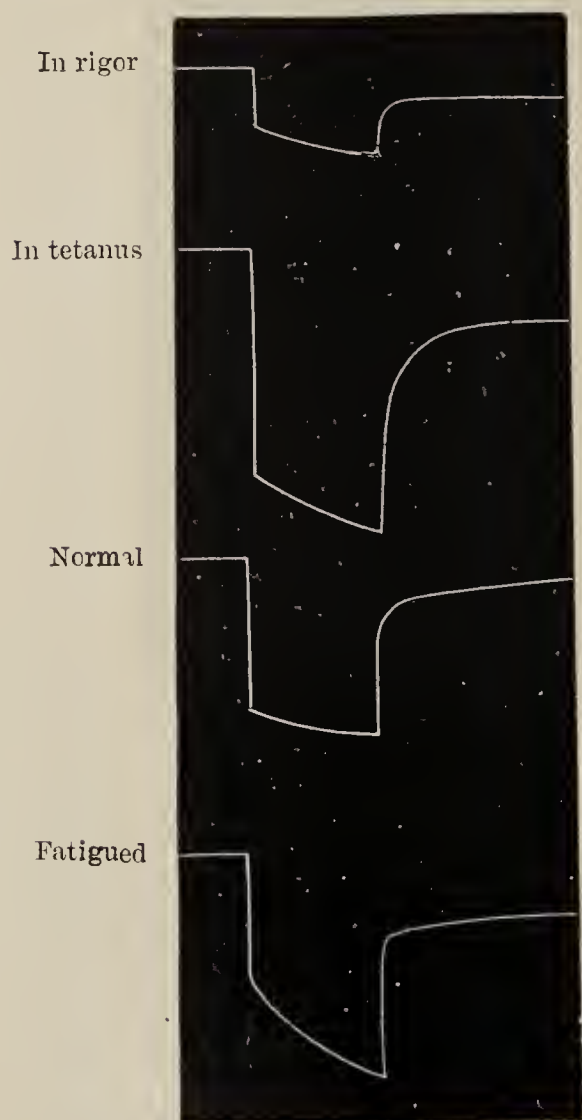


FIG. 137.—EXTENSIBILITY OF MUSCLE IN VARIOUS STATES.

Tested by 50 grammes applied for short periods.

one, and that the recovery is not quite perfect. It is, however, usual, and no doubt correct, to say that the elasticity of muscle in its normal relations is perfect; though, out of the body and abnormally stretched, the muscle does not exhibit this perfect elasticity. Active muscle is more extensible than resting muscle; rigid muscle is less extensible than normal muscle, and its elasticity is very imperfect.

If a muscle be weighted and left so, it will be noticed that the weight causes an immediate elongation, followed by a gradual elongation, which continues in a diminishing degree for an indefinite time. If a weighted muscle be relieved of weight, it will be noticed that the immediate elastic shortening is followed by a gradual indefinite shortening.

These after-effects (viscosity) are not characteristic of muscle, but common to all extensible and elastic substances; they are best demonstrated by causing the

recording lever to mark its excursion upon a slowly-travelling surface, *e.g.* a smoked cylinder fixed to the hour-axis of an ordinary clock.

Muscular extensibility and elasticity are useful in modifying sudden muscular contractions in the living body, rendering the application of their force more gradual, and thus obviating sudden jerks or rupture of tissue; moreover, the muscles are normally slightly stretched between their points of attachment to the bones, and are thus favourably disposed for prompt commencement and smooth execution of movements.

**Contractility.**—In common with all living tissues, muscle possesses excitability, which property manifests itself as shortening

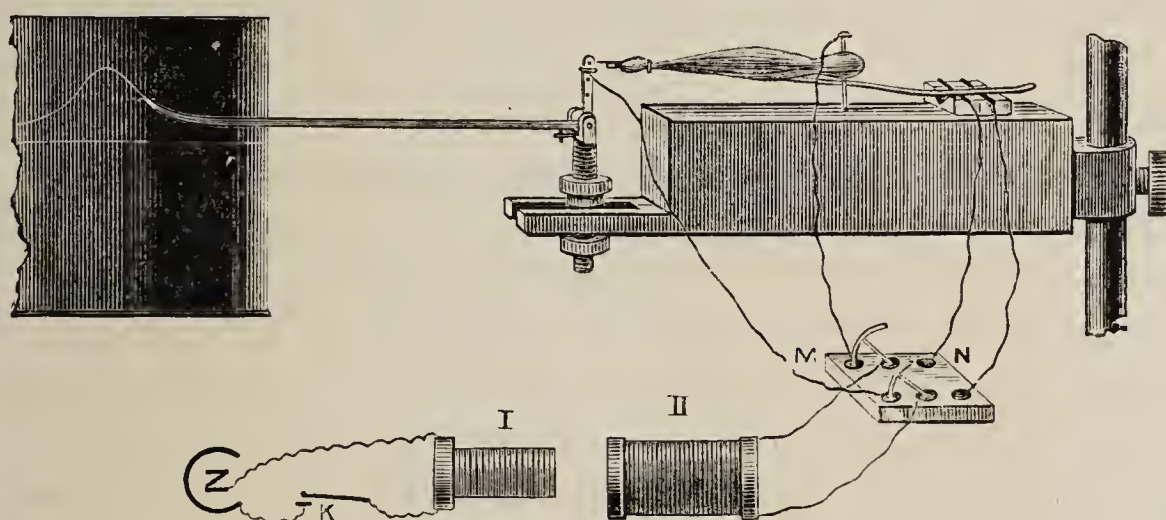


FIG. 138.

Sketch to show a myograph and commutator without cross-wires arranged for direct and indirect excitation. Wires from secondary coil to middle pools of commutator; fine wires from left-hand pair of pools to tendon of muscle and to pin through femur (circuit M), and from right-hand pair of pools to a pair of electrodes across which the nerve is laid (circuit N); moving the cradle to left or right connects the secondary coil with muscle or with nerve, and current is sent through either by opening and shutting a key K in the primary circuit.

or contraction; we say, therefore, that living muscle is *contractile*, *i.e.* able to contract, and that its characteristic vital property is '*contractility*,' *i.e.* ability to contract. Any agent that excites a muscle to contract is called a '*stimulus*'; it may be mechanical, thermic, chemical, electrical, or physiological. Of these various kinds of stimuli, that which is usually employed experimentally is the electrical stimulus, and the electrical stimuli generally used are (1) the constant current and (2) induced currents. Normal muscular movement in the body is excited by the physiological stimulus which rises in the brain and is conveyed by motor nerves to muscle. Mechanical, thermic, and chemical stimuli are but little employed to stimulate muscle directly. The application of an experimental stimulus directly to muscle constitutes *direct*



stimulation; the application of a stimulus to its motor nerve constitutes *indirect* stimulation, these expressions being used with especial reference to '*nerve-muscle preparations*,' which are usually composed of a sciatic nerve in connection with a gastrocnemius muscle removed from a recently killed frog.

**Constant current.**—Stimulation with the constant current is effected when the current of a battery begins to pass and when it ceases to pass through a muscle, *i.e.* when it is *made*, and when it is *broken*, by closing and opening a key in the circuit of which a nerve or a muscle forms part. The muscle will contract each time the key is closed (make or closure contraction) or opened (break or opening contraction); it will remain quiescent while the current passes, *i.e.* while the key is left closed.

This is the rule to which, however, there are exceptions; what is known as '*Wundt's tetanus*' is an enduring contraction that is apt to occur in a frog's muscle after injury or during the passage of a strong current; and on human muscle the effect termed '*galvano-tonus*'—which occurs on normal muscle during the passage of a strong current, on degenerating muscle during that of a comparatively weak current—is of a similar character.

If the stimulus is applied at one end of a muscle, the proximal portion begins to contract a little earlier than the distal portion, *i.e.* the contraction spreads as a wave from the point of excitation, its rate of propagation (in frog's muscle) being between 1 and 3 meters per second (Aeby).

In perfectly normal fresh muscle the contraction sweeps undiminished from end to end; in moribund muscle it is stronger at beginning than at end, or it may actually remain localised to the point excited; this is the '*idio-muscular contraction*'; on weakly subjects it may be easily elicited by mechanical excitation.

If a strong galvanic current is passed longitudinally through a freshly-excised and gently-extended sartorius muscle of a frog, a series of undulations may be seen to sweep along the muscle in the direction of the current (*i.e.* from anode to kathode) with a velocity of 3 to 5 mm. per second (Kühne, Hermann). The effect, which is not to be confused with the normal wave of contraction, is an idio-muscular phenomenon, probably caused by intrapolar irregularities of current-density; moderate warmth is favourable to its production.

Comparing the effects at make and break of the constant current, it will be found that the contraction at make is stronger

than the contraction at break ; to determine this relation it is necessary to graduate the strength of current and consequent strength of stimulation by means of the rheochord (p. 305). It will be found by further experiments that *the make contraction starts from the kathode, that the break contraction starts from the anode* ; in other words, that *the make stimulus is kathodic, the break stimulus anodic*. Experiments demonstrating this statement are as follows :

(I.) Two levers rest upon a curarised muscle near its two ends, to which the kathode and anode respectively are connected, and are arranged against a recording surface ; on making the current both levers rise, the kathodic a little sooner than the anodic ; on breaking the current both levers rise, the anodic a little sooner than the kathodic.

(II.) A muscle is injured at one end and stimulated by make and break of a constant current, first in one, then in the opposite direction. It is found that when the kathode is at the injured end the make stimulus is less effectual than on the uninjured muscle ; when the anode is at the injured end, the break stimulus is less effectual than on the uninjured muscle.

The statement made above applies to smooth as well as to striped muscle, although an apparent exception usually occurs in the former, and sometimes in the latter case, *i.e.* a contraction apparently started at the anode as well as at the kathode ; on closer examination it is, however, found that whereas excitation starts from the precise point with which the kathode is in contact, it starts from an area at some distance surrounding the anode (Biedermann). We shall encounter a similar phenomenon and shall give its explanation in dealing with the excitation of human nerve.

In the undifferentiated protoplasm of protozoa, the polar response appears to be opposed to that obtaining in differentiated muscle (and nerve). Whereas in the latter the response is to kathodic make and anodic break, in unfibrillated protoplasm it is to anodic make and kathodic break.

One of the most remarkable reactions exhibited by these lowest forms of animal life is *galvanotropism* ;<sup>1</sup> many of the infusoria of *e.g.* a drop of ' hay infusion ' traversed by a galvanic current travel in the lines of current towards the anode or the kathode. Ciliata travel with the current and collect at the kathode ; flagellata travel against the current and collect at the anode (Verworn).

<sup>1</sup> τρέπω or τρωπάω, to turn.



Animals higher in the scale, such as young tadpoles and fish, also exhibit galvanotropism, which, however, appears to be a spinal reflex reaction; if they are placed in a trough of water traversed by a strong galvanic current they adjust themselves longitudinally to the lines of current, with heads to the anode, turning round as soon as the direction of current is reversed (Hermann).

**Induced currents.**—In the application of induced currents there is no direct connection of the battery with the muscle—the latter is connected with the *secondary* coil, and is stimulated by currents induced in that coil when the battery current begins and ceases to flow through the *primary* coil. The current that is induced in the secondary coil when the battery current begins to flow through the primary coil, is the make induced current; that which is induced in the secondary coil when the battery current ceases to flow, is the break induced current. These two currents are in opposite directions, and are not to be confused with the make and break of a constant current. They are of momentary duration, the beginning and end of each induced current being practically simultaneous, so that the stimulus of each is compounded of an instantaneous rise and fall of current.

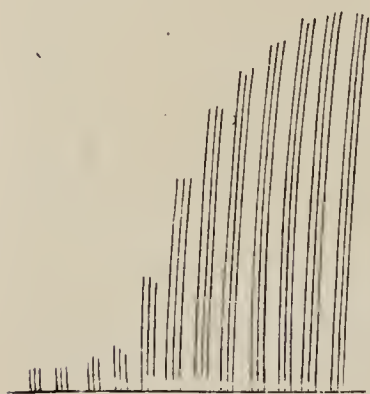


FIG. 139.

Increasing height of muscular contraction with arithmetically increasing strength of stimulation; at each strength a group of 3 isotonic contractions is recorded.

Induced currents are the most frequently employed test of excitability; their strength is varied by varying the distance of the secondary from the primary coil; in the case of muscle, for instance, excitability is determined by finding the greatest distance of secondary from primary at which response is obtained, or by observing increase or diminution in the height of contractions obtained, while the two coils remain at a distance previously determined to give a moderate strength of stimulation. Comparing the effects at make and break of the primary current, *i.e.* the effects of the make induced current and of the break induced current, it will be

found that the contraction caused by the break induced current is stronger than that caused by the make induced current, or, what amounts to the same thing, that the contraction at break will be produced by a weaker current than the contraction at make. This is not a specific difference comparable with the differences between contractions provoked by make and break of a constant current, but simply an evidence

of the fact that the break induction current is sharper than the make induction current. The two contractions can be equalised by modifying the connections of the primary coil according to the plan first used by Helmholtz (see p. 315).

A '*maximal*' stimulus is one that causes a muscle to perform the greatest contraction of which it is capable as the effect of a single stimulus; a stimulus of greater strength cannot therefore give rise to a greater effect. A '*submaximal*' stimulus is one that causes a muscle to perform a contraction smaller than that which is brought about by a maximal stimulus. A '*minimal*' stimulus is the weakest stimulus that will cause any contraction at all.

With regard to the relation between the magnitudes of stimulus and of contraction within the range from minimal to maximal, it appears that the lift increases with an at first increasing and subsequently a diminishing increment, so that its curve of increase is at first convex then concave towards the abscissa, viz. S-shaped.

**Myographs.**—Each induction current gives a single muscular contraction or twitch, the form of which is studied by the graphic method. A lever in connection with a muscle, and rubbing against a travelling smoked surface so as to give a magnified record of the movements of the muscle, constitutes a *myograph*. Myographs are of many kinds and shapes, according to special requirements. The recording surface may consist of a cylinder made to revolve by clockwork; or of a plate on a carrier which is shot along horizontal guides by a spring; or of a plate fixed to a pendulum. In every case the rate of movement of the surface is to be determined by a *chronograph* marking time simultaneously with the movements of the muscle-lever. A vibrating tuning-fork with a pen fixed to one of its prongs, or a vibrating reed interrupting the

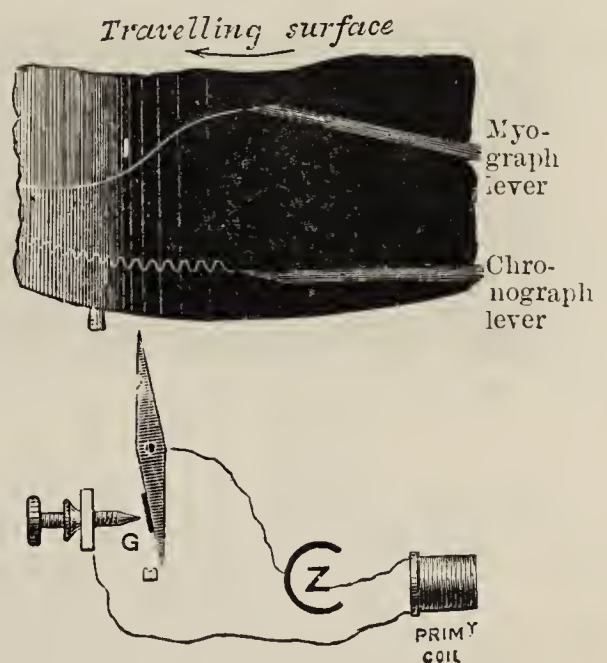


FIG. 140.

Fixed key opened by a recording cylinder, as used for taking a latent period. The point of stimulation is marked by touching the myograph lever with the peg in contact with the shut key; the lost time is ascertained from the interval between that point and the rise of the curve described when the recording surface has passed at full speed and struck open the key.



circuit of a small electro-magnet with a pen fixed to its armature, constitutes the chronographs in most frequent use. A tuning-fork vibrating 100 times per second traces a series of teeth, each of which indicates  $\frac{1}{100}$ th second; similarly, a reed vibrating 20, 50, 100, or 200 times per second, interrupting a current an equal number of times, traces the time-indications in  $\frac{1}{20}$ ths,  $\frac{1}{50}$ ths,  $\frac{1}{100}$ ths, or  $\frac{1}{200}$ ths.

*A simple contraction or twitch.*—A single induction shock (make or break induced currents) led into a muscle causes a single contraction or twitch. If, instead of producing the make or break in the primary circuit by a key moved by hand, this be done by a key that is closed or opened by a peg fastened to the travelling surface (cylinder or plate), it will be easy to observe that the muscle does not contract at the precise instant of stimulation, but a fraction of a second later. This interval between stimulation and effect is called the '*latent period*,' or physiological lost time.

The length of the latent period in voluntary muscle is  $\frac{1}{100}$ th second (Helmholtz),  $\frac{1}{200}$ th second (Yeo),  $\frac{1}{400}$ th second (B. San-

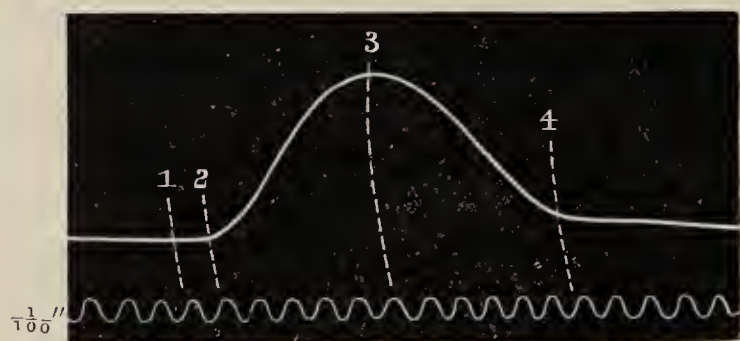


FIG. 141.—A SINGLE MUSCULAR CONTRACTION  
(Frog's Gastrocnemius).

From 1 to 2 is the latent period; from 2 to 3, the period of shortening; from 3 to 4, the period of relaxation.

The last-named figure is the shortest obtainable under the most favourable conditions and is not perceptibly longer than the latent period of the current of action. There is, in fact, reason to believe that the contractile elements of the muscle are so inobedient that there is no appreciable interval between a

direct stimulus and the commencement of their mechanical response. And the appearance of a mechanical latency in the entire muscle subsequent to the manifestation of an electrical variation can be accounted for by the fact that it takes time for the contractile effect to spread along the whole muscle; owing to the extensibility of the muscle, the part at which contraction starts must first stretch the rest of the muscle, and a lever suspended from it will not begin to rise until the total active shortening has exceeded the total passive elongation.

If a maximal stimulus be applied—(a) directly to muscle, (b) to the nerve close to its entrance—the lost time of the second

contraction is found to be  $\cdot 005$  sec. longer than that of the first, a delay that is attributable to the interposition of the motor end-plates.

The latent period is followed by the muscular contraction which, when recorded as just described, consists of an ascending portion, during which the muscle shortens to a maximum, and a descending portion, during which the muscle returns to its original length. The total duration of a single muscular twitch is about  $\frac{1}{10}$ th second.

A muscle to which a weight is suspended is said to be '*loaded*'; if a support is so adjusted that the muscle raises the weight only during contraction, and is not stretched by the weight when at rest, the muscle is said to be '*after-loaded*.' If a muscle contracts against a small and constant resistance, so as to be extended by a constant force during its contraction, the curve described by a light lever attached to it is termed '*isotonic*.' If a muscle contracts against a large resistance, *e.g.* a strong spring, so that it can shorten very little, the curve described by a lever attached to it is termed '*isometric*.' The latter is flat-topped, *i.e.* exhibits a period of maintenance at maximum contraction. As ordinarily employed, the myograph gives an isotonic curve; the isometric method is exemplified by the spring '*dynamometers*' used for clinical purposes.

*Summation.*—If a muscle is made to receive two stimuli in rapid succession, the effects of each will be added together and produce a greater effect than either separately; the second contraction will be superposed upon the first. This is a summation of *effects*, as distinguished from summation of *stimuli*.

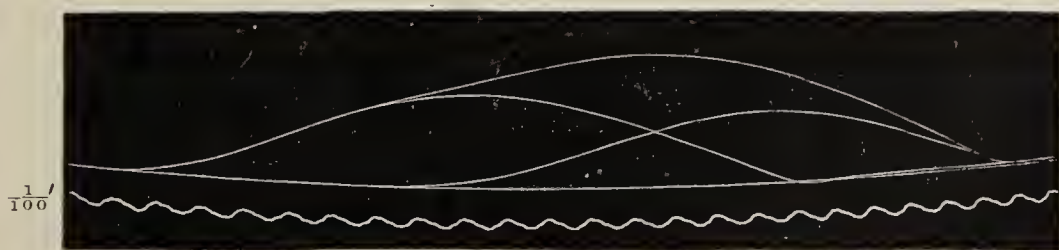


FIG. 142.—SUPERPOSITION OF TWO SINGLE CONTRACTIONS.

Each contraction is recorded alone by a break shock caused by opening a fixed key; both keys are then set, and the recording plate striking them open successively, causes two stimuli and a summation of the two contractions.

If the two stimuli are in such close succession that the second occurs during the latent period of the first, the result will differ according as the stimuli are maximal or submaximal. If they are maximal, the contraction will be produced by the first stimulus, and will be unaffected by the second stimulus; if they are



submaximal, the two stimuli will become added together, and produce a greater effect than either separately. This is summation of stimuli.

A succession of stimuli will cause a succession of contractions, the number of contractions corresponding with the number of stimuli. As the frequency of these is increased, so the frequency of the individual contractions increases. The muscle now does not completely relax, but remains more or less permanently contracted, so that the point of the lever attached to it is kept above the level at which it would trace if the muscle were relaxed, and makes from this new level smaller excursions corresponding with individual contractions. This is incipient or incomplete tetanus,

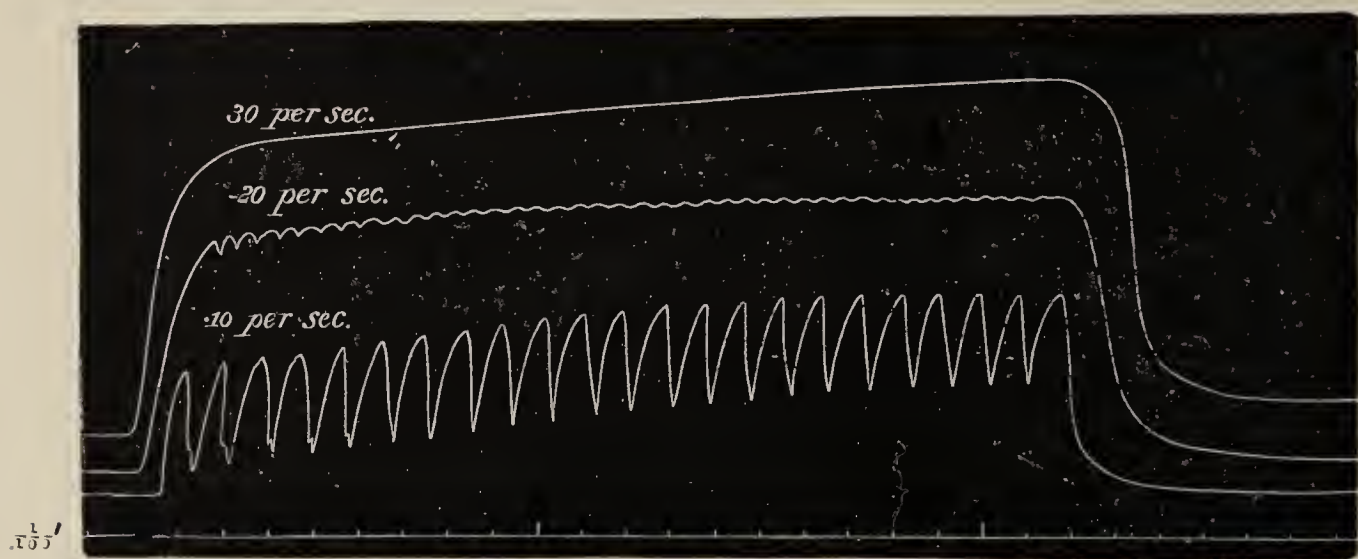


FIG. 143.—COMPOSITION OF TETANUS.

Stimuli caused by a spring interrupting a primary circuit by vibrating in and out of a mercury cup; the vibration-frequency is increased by shortening the spring (see p. 317).

or, as it is sometimes called, '*clonus*.' With still further increase in the frequency of stimulation, the individual contractions become completely fused, the muscle remains in complete contraction so long as the stimulation is kept up, and the lever traces an unbroken line during this period. This is complete *tetanus*. The smallest frequency of stimuli required to produce it varies between 20 and 50 per second for ordinary striated muscle, according to temperature and fatigue. Beyond this minimum any further increase of frequency (strength and quickness of stimuli remaining constant) gives no further increase in the amount of muscular shortening.

Alternating currents (*i.e.* currents oscillating to and fro in opposite directions) of high vibration-frequency, and strong enough to heat a carbon filament to incandescence, can be passed through muscle (or nerve) without producing excitation (Tesla,

d'Arsonval) ; whereas a constant current of  $\cdot 005$  ampère through the human body is excitatory both at make and break, a current of 5 ampères (?) but oscillating at a vibration-frequency of 1,000,000 per second, is unfelt and produces no contraction ; with suitable apparatus we may send such a current through one or more persons and electric lamps in the same circuit, and see the lamps illuminated by a current that is not felt.

The contraction of striated muscle varies in different animals and in different parts of the same animal, and may be modified by fatigue, by temperature, and by drugs. In different animals some muscles subserve short sharp movements, others prolonged movements ; their individual contractions are correspondingly short or long ; *e.g.* in the crayfish the muscles that move the tail give short twitches, those that move the claw give long grasps ; in the frog, the gastrocnemius gives a longer contraction than the hypoglossus ; in the rabbit, the soleus (red muscle) gives a longer contraction than the gastrocnemius (pale muscle) ; in man, the muscles of the upper extremity give shorter contractions than those of the lower extremity.

A muscle that from any cause gives a long contraction is tetanised at a lower stimulation-frequency than a muscle giving a short contraction. Red muscle, cooled muscle, fatigued muscle, have a lower minimum tetanising-frequency than pale muscle or normal muscle. Rigor comes on later and lasts longer in red than in pale muscle.

*Fatigue* affects muscular activity, causing a diminution of absolute force and of available work, and a general sluggishness of movement ; it is attributable to two causes conjointly—viz. to the consumption of force-producing material, and to the accumulation of force-impeding material. The record of a series of muscular contractions exhibits the gradual development of the slowing and declining activity, after a short period of exaltation. The entire process of fatigue is divisible into two stages : (1) a short preliminary stage, during which the contractions increase in height and in duration ; (2) a longer stage, during which they continue to increase in duration but progressively diminish in height. Figs. 144 and 145 illustrate these points, and fig. 146, given to show the effects of diminishing temperature, might equally well be an example of the two stages in a fatigue effect. The curve of decline in the (isotonic) contractions of an after-loaded frog's muscle stimulated at regular intervals is a straight line (Kronecker). With a series of isometric contractions the fatigue



curve is S-shaped, *i.e.* at first concave, subsequently convex towards its abscissa. (w.)

If left to itself, a muscle that has been exhausted—*i.e.* forced into action until it can act no longer—will recover; normally, in the body a muscle is preserved from exhaustion, and even from

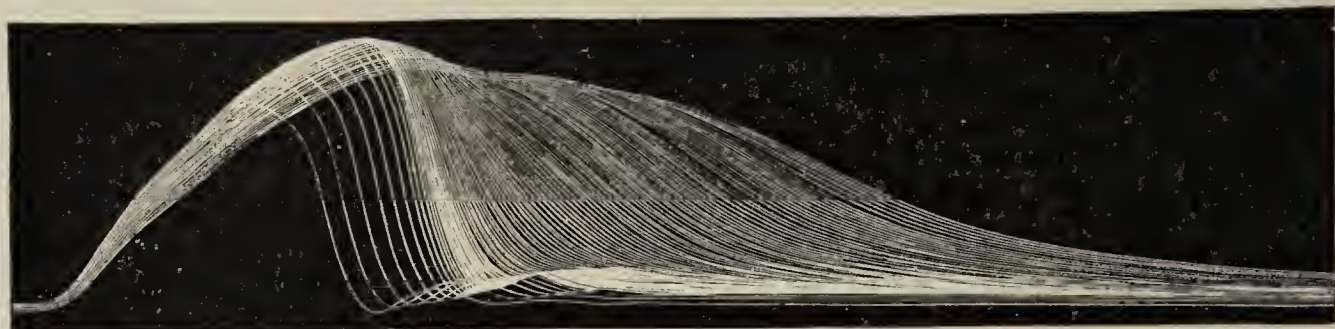


FIG. 144.—FATIGUE. (Frog's Gastrocnemius.)

Direct excitation; 125 successive maximal contractions at intervals of  $1\frac{1}{2}$  sec., showing at the outset increase of height and of duration, later decreasing height. (The exhaustion has not been pushed to the end.)

appreciable fatigue, by the circulating blood bringing fresh matter and carrying off fatigue products. We shall see later (p. 383) that the exhaustion of muscle by nerve action is rendered impossible by the comparative instability of the end-plates, but the point which we should now clearly recognise is, that even an *excised bloodless*

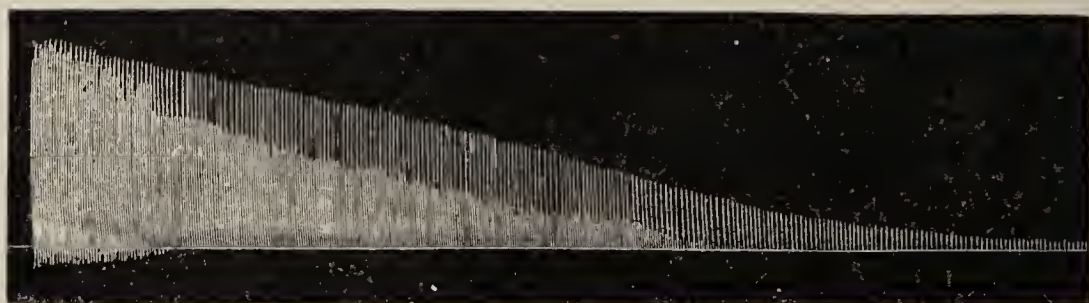


FIG. 145.—FATIGUE. (Frog's Gastrocnemius; isotonic contractions.)

About 250 make and break stimuli. (The recording surface travels very slowly, so that each contraction marks a vertical line only; the make disappear sooner than the break effects.)

*muscle* has within itself material and power of self-restoration after exhaustion, independently of new material furnished by the blood; left to itself, an isolated bloodless muscle recovers contractility.

A given muscle contracts more or less suddenly and sharply according as *temperature* is high or low. As the effect of cold the record of a single contraction exhibits (1) a longer latent period, a prolonged and a higher contraction, (2) a still longer latent period and a still more prolonged, but lower, contraction. Thus, as might be expected, the effects of lowered temperature are identical

with those of fatigue, and in both cases are attributable to greater sluggishness of change. As the effect of increased temperature, each contraction becomes shorter and smaller; the excitability, after a slight increase, diminishes and is lost; finally, when the

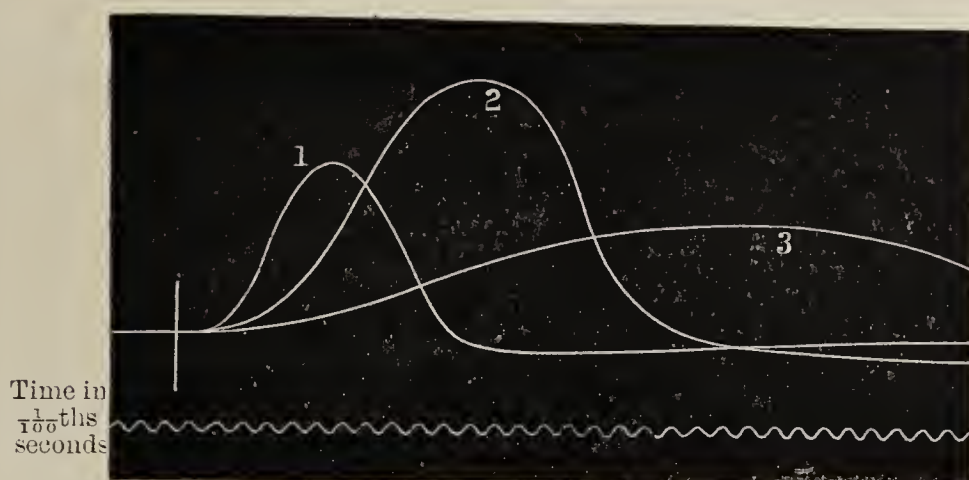


FIG. 146.—EFFECT OF TEMPERATURE UPON MUSCULAR CONTRACTION.

1. Normal. 2. Cooling. 3. Very cold.

temperature exceeds  $40^{\circ}$ – $50^{\circ}$  C. the muscle shortens and becomes rigid (heat-rigor). In this rigor the shortening exceeds that produced by a maximal tetanus, but the absolute force of the contraction is very small.

A saline extract of exhausted muscle depresses the excitability of fresh muscle through the vessels of which it is made to circulate (Ranke). The blood of a dog exhausted by excessive exercise causes ‘symptoms of fatigue’ if transfused into the vascular system of a fresh dog (Mosso). These effects are attributable to ‘fatigue products,’ lactic acid, and other bodies unnamed.

Among drugs causing a modification of contraction by their direct action upon muscle, the most notable is *veratrin*, which, if

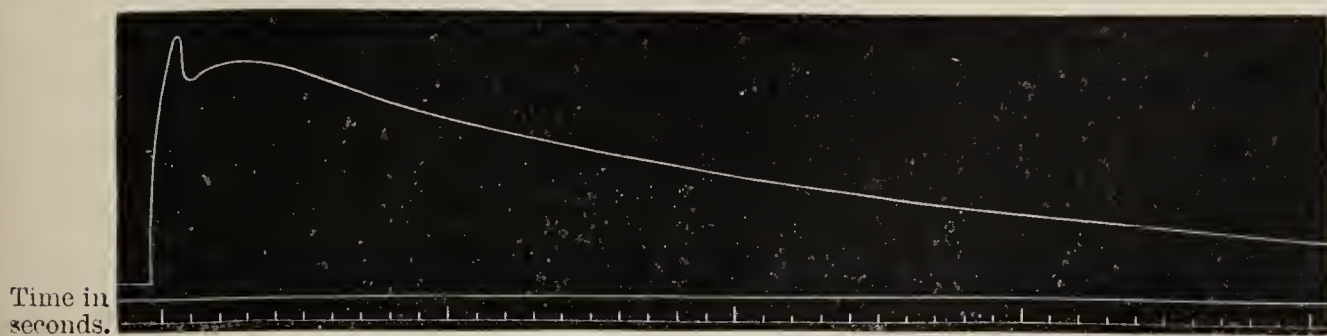


FIG. 147.—VERATRIN CURVE.

injected into an entire frog, or directly applied to an excised muscle, causes an excessive prolongation of the muscular contraction. Among inorganic bodies, the *calcium* salts may be mentioned as producing a similar though less marked effect.



*Proteid solutions* nourish or fail to nourish skeletal muscle, just as they nourish or fail to nourish cardiac muscle; serum-albumin is 'nutritive,' egg-albumin and the digestive proteids (albumose and peptone) are non-nutritive, and the latter are actually 'poisonous' (J. Brinck).

**The muscular sound. Voluntary contraction.**—Contracting muscle emits an audible sound; with a single contraction a short thud is heard; during a tetanic contraction a purring or rumbling noise is heard, having a pitch of about forty per second. But, as we shall see, this tone is no proof that impulses are generated with corresponding frequency, the pitch heard being in reality dependent upon the resonance tone of the listening ear. Upon muscles tetanised by electrical stimulation, of, *e.g.*, 50 to 100 per sec., tones are heard corresponding in pitch with the stimulation-frequency; and it is stated that in muscular tetanus produced by high stimulation-frequencies (1,000 Bernstein, 700 Lovèn), a note of corresponding pitch, or of the next octave below, is audible; in observations of this kind it is necessary to guard against a fallacy due to the static effects of alternate charge and discharge; these, if they should occur, will cause an audible tone corresponding with the interruption-frequency, but independent of any muscular contraction, and equally well heard on dead muscle (Lovèn). The fallacy is obviated by touching a moist surface of the animal with a moist finger during the auscultation.

Every voluntary contraction, even the shortest, is considered to be a tetanic series of impulses; nevertheless, it is possible to make a single voluntary contraction as short as a single induction twitch, or to cut it even shorter by the antagonist contraction. Wollaston, who first studied voluntary muscular contraction by the acoustic method, came to the conclusion that the muscle-note emitted during voluntary contraction has a vibration-frequency of 36 per second. Helmholtz, by means of consonating springs, concluded that the actual number of neuro-muscular impulses producing this effect is 18 per second; and he pointed out as a fallacy of the acoustic method that the pitch of the sound depends, not upon the observed muscle, but upon the observer's ear, of which it is a resonance tone; moreover, the tone of a muscle in galvanotonic contraction, *i.e.* in the continuous contraction produced by a *strong* continuous current, has the same pitch as that of a voluntarily contracted muscle (w.) Lovèn called attention to the fact that strychnia spasms in the frog have

a rhythm of 8 to 10 per second; and considers this to be the normal rate of spinal discharge.

The maximum number of *double* (*i.e.* to and fro) voluntary movements which we can make in one second is 8 to 10; the smallest interval at which two separate motor impulses can be made to succeed each other is  $\cdot 10$  to  $\cdot 12$ ; the rhythm of morbid spasms—trepidation, clonus, tremor—is 8 to 10 per second, so also is that of clonic tremors occurring in normal muscle, especially in the condition of fatigue or of shivering (*w.*); Schäfer and v. Kries have independently pointed out that the graphic record of even the steadiest voluntary movement exhibits a tremor of about the same frequency (8 to 12), which—if it was not due to vibrations of the recording levers (Haycraft)—must have been a similar phenomenon in its most normal and rudimentary form. The sum of these various items signifies to us that the physiological mobility of the central nervous system is characterised by the maximum impulse-frequency of about 10 per second.

*Is the heart-beat a long twitch or a short tetanus?* We are now in a position to answer this question, and the state of our knowledge is such that the answer can be placed on surer ground than at a time when it rested upon interpretations of secondary contraction and of tetanic bruits. We need only very briefly consider these two data. (1) When the nerve of a nerve-muscle preparation is laid across the beating heart, the muscle gives a single twitch, at the outset of each beat, and occasionally also at its close; this was received as an indication that the heart's contraction is single, and not multiple, and we shall find in a further section that this evidence is much strengthened by our present knowledge of its meaning; it is an index of the diphasic variation that accompanies each single contraction (p. 390). (2) The sound made by the contracting heart was formerly much quoted in support of the opposite view, according to which the systole was considered to be a short tetanus. The argument was as follows:—muscular sounds are due to a tetanic succession of impulses, the heart gives a sound during its contraction, therefore that contraction is tetanic. But we now admit that any contraction, even a single twitch, gives a sound, short if the contraction is short, long if the contraction is long; the argument falls, therefore, to the ground. We can no longer hesitate to admit that each contraction of the ventricles is a single spasm; comparing the contractions of striated muscle, of cardiac



muscle, and of smooth muscle, we recognise that the only difference is one of duration; the latent period, the rise and fall of contraction, maintain their relation to each other, and—in the cases of skeletal and of cardiac muscle—the passage of a single wave of change has been followed from origin to end along the

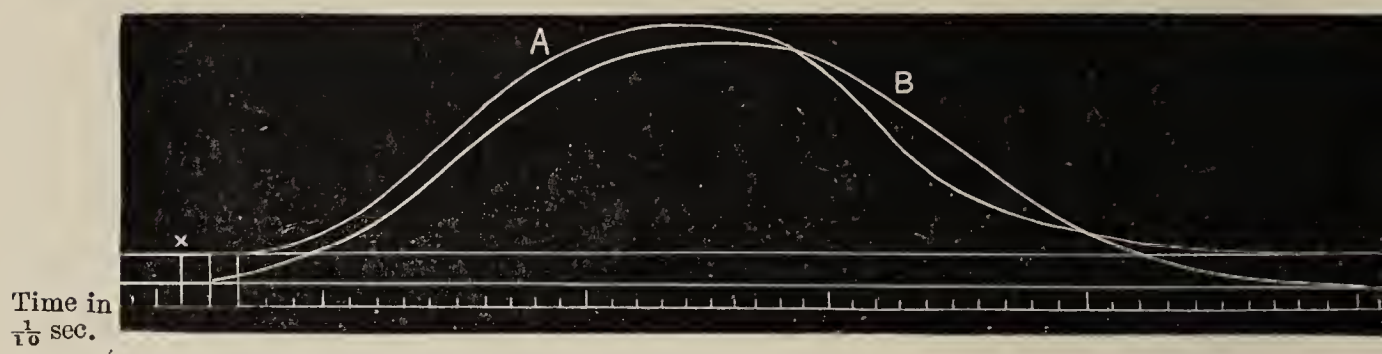


FIG. 148.—WAVE OF CONTRACTION IN FROG'S VENTRICLE, TAKEN BY TWO LEVERS RESTING UPON IT NEAR THE APEX AND NEAR THE BASE.

x marks the moment of excitation, which is applied near the base, *i.e.* in imitation of the normal origin of the contraction. The contraction of the base begins sooner and lasts longer than that of the apex.

excitable tissue—electrically, as we shall see on p. 390, and also mechanically, as is illustrated above.

**Strength; work.**—The *absolute strength* of a muscle is measured by the weight that a muscle just fails to lift on being stimulated, the weight being applied as an 'after-load.' This weight varies with the sectional area of muscle, not with its length; *i.e.* the absolute strength of a thick muscle is greater than that of a thin muscle; and the absolute strength of a long muscle is the same as that of a short muscle, if both are of the same thickness. The absolute strength of tetanised frog's muscle has been found to be about 3 kilogrammes per 1 square centimeter sectional area; of human muscle it is greater, viz. 5 to 10 kilogrammes per 1 square centimeter. This is for voluntary contraction; the strongest possible artificial tetanus only reaches  $\frac{1}{2}$  to  $\frac{2}{3}$  of the voluntary maximum. The absolute strength in tetanus is about double that of a single contraction in the frog, and no less than ten times its value in human muscle (Fick).

The *work done* by a muscle is measured by the product of weight raised  $\times$  height. A muscle that shortens 10 millimeters, raising a weight of 100 grammes, does work equal to 1,000 gramme-millimeters, or 1 grammeter. Leaving out of account the weight of the muscle itself, no work is done when muscle contracts without raising a weight or overcoming some resistance. A tetanised muscle does work only at the beginning of tetanus, when it raises a weight; during the tetanus, while the weight is

kept up, the contraction is static, and no work is done. If we measure the work done by a muscle made to raise a succession of increasing weights, we shall find that it increases to a maximum, from which it declines with further increase of weight.

Work done thus depends upon three factors—(1) amount of weight, whether as ‘load’ or as ‘after-load’; (2) length of muscle; (3) sectional area of muscle. Length  $\times$  sectional area is equivalent to bulk or weight; the maximum work that can be done is therefore dependent upon the bulk or weight of the muscle in action. It has been found that for frog’s muscle this maximum under favourable circumstances is in a simple contraction nearly 1 gram-meter per 1 gramme, and in a short tetanus about 4 grammeters per 1 gramme.

An ordinary labourer at work does about  $\frac{1}{5}$  gram-meter per gramme muscle per second; *i.e.*, taking his muscle to be 20 to 25 kilogrammes, his work during 8 hours = 115,000 to 144,000 kilogrammeters.<sup>1</sup> A man during an ordinary walk does work at about the same rate. A long-distance bicycle-racer was calculated to do work for 8 hours at the rate of 1 grammeter per gramme per second, the weight of his muscle being estimated at 25 kilogrammes; a rowing man was calculated to work at the same rate, but only for a short time; the work of a short-distance runner has been estimated at between 2 and 2.5 grammeters per gramme per second. According to Marey, the work done in walking and running on level ground amounts to between 10 and 20 kilogrammeters per step.

The muscular strength is tested on man by means of the *dynamometer*, which in its ordinary clinical form consists of a strong oval spring to be grasped in the hand, the value of the

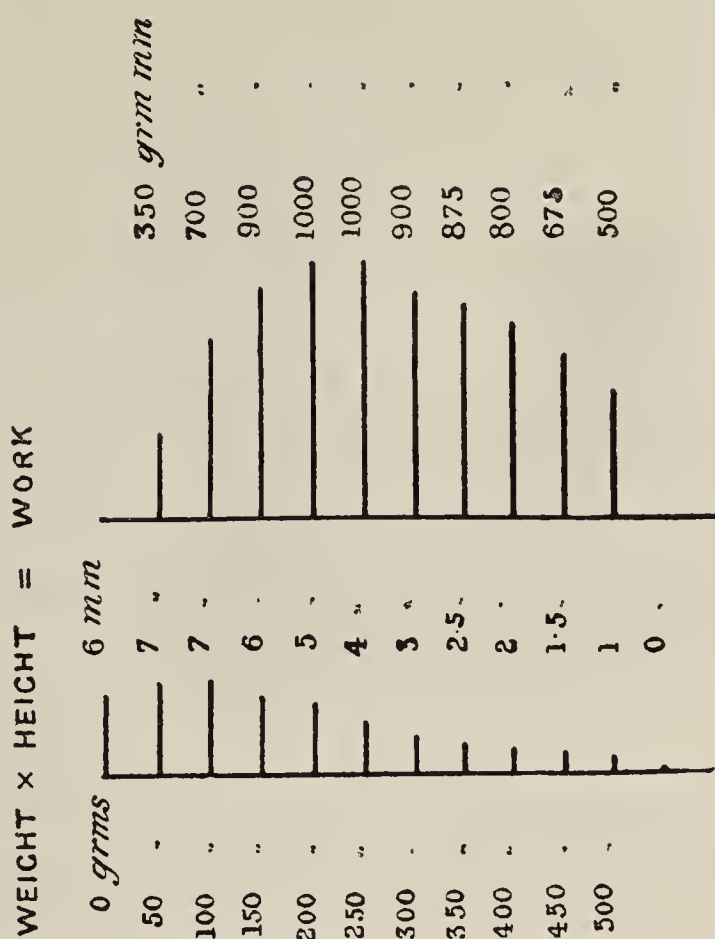


FIG. 149.

<sup>1</sup> This is a low estimate; 1 horse-power = 77 KgM. per sec.; 1 man-power =  $\frac{1}{8}$ th horse-power, or nearly 10 KgM. per sec., *i.e.* 288,000 KgM. per 8 hours.



flattening being shown by an index and scale graduated in pounds or kilogrammes. The value of the instrument is much diminished by the fact that habit and 'knack' are very important factors in the maximum squeeze: its indications are of greater value if the instrument is converted into a *dynamograph*; the manner of maintenance of a prolonged maximum effort for regular periods, or the character of a series of maximum efforts for regular periods, may then be recorded on a slowly-travelling surface, *e.g.* a cylinder on the hour-axis of an ordinary clock; and from such records an estimate may be formed of the muscular strength, of its rate of decline in a succession of efforts, and of its rate of recovery from such decline (w.)

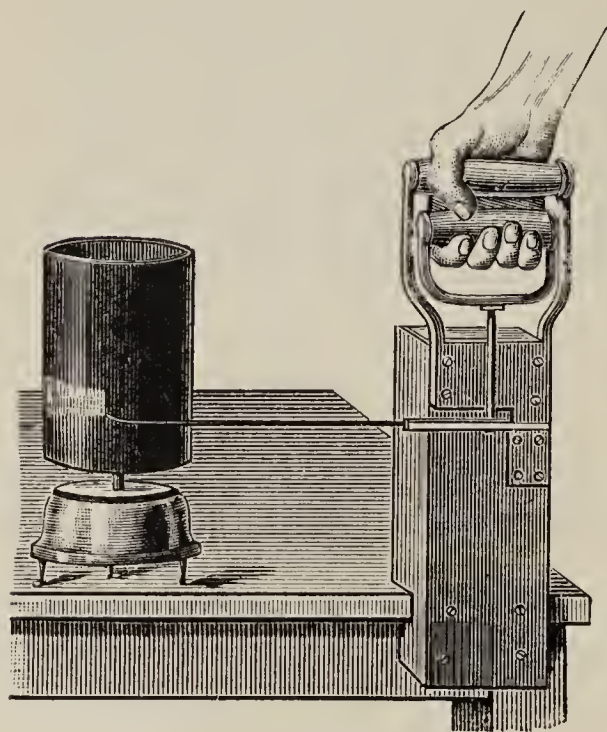


FIG. 150.—DYNAMOGRAPH.

Short straight spring and long lever (represented here much shorter than it is in reality).

of work as kilogrammeters, although in other respects the isometric method is preferable. The half-supinated arm is attached to a horizontal support, a cord from a ring round the

By means of an instrument devised by Mosso, and termed by him the *ergograph*, similar estimates may be made, with the advantage that they may be numerically expressed in terms

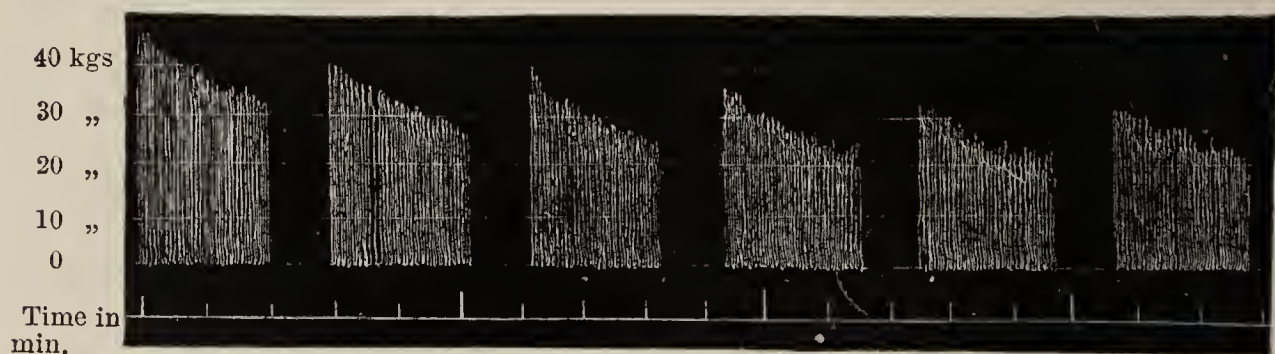


FIG. 151.—DYNAMOGRAPH TRACING.

Each group of lines is the effect of 30 maximal efforts, each lasting two seconds. Intervals of rest for one minute between successive groups.

middle finger passes over a pulley and carries a weight, the hand is kept in position by two tubes into which the index and ring fingers are inserted; the successive elevations of the weight by flexion of the middle finger are recorded in the usual manner.

**Heat evolved.**—The chemical action taking place in muscle is

of course accompanied with an evolution of heat, and its amount is increased during contraction. It has been found that, *cæteris paribus*, (1) more heat is evolved in the contraction of a stretched than in that of an unstretched muscle, (2) more heat is evolved by a muscle that contracts without doing work than by a muscle that does work (Heidenhain). These facts have a far deeper significance than is superficially apparent. The first fact implies that pure muscle is not a mere machine discharging energy irrespectively of the work to be done, but that it evolves much or little energy according as it pulls against much or little resistance, and it is obvious that this relation between demand by tension and response by action is an important feature in the economy of natural contraction. The second fact is an illustration of the conservation of energy: if the total energy set free appears in the form of heat, there is more heat than if some of the energy goes off as work. The most favourable conditions for the demonstration of the heat of muscular contraction are therefore established by tetanising stretched muscle which cannot shorten and do work.

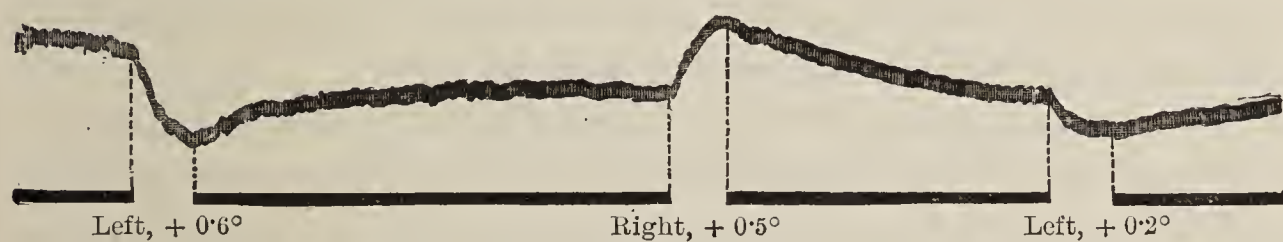


FIG. 152.

Two thermo-electric needles in the two gastrocnemii of a frog, and connected with a low-resistance galvanometer; the spot of light from the mirror records the deflection of the magnet upon a cylinder covered by sensitive paper and revolving round a horizontal axis. Excitation of the sciatic on one side for 1 min. causes increased heat in one muscle and deflection of the magnet; excitation on the opposite side gives deflection in the opposite direction. Periods of excitation indicated by breaks of the abscissa. In this instance the rise of temperature is considerably greater than that generally quoted, viz., according to Helmholtz, a rise of  $0.1^{\circ}$ , with a tetanus lasting one minute; according to Fick, a rise of  $0.002^{\circ}$  with a single contraction.

Considered in a slightly different manner, these two important facts become still more intelligible. In the case of the first, which may be designated by the aphorism '*Ut tensio sic calor*,' we must be careful not to suppose that a muscle raising a heavier weight, *i.e.* under greater tension and doing more work, yields this mechanical energy at the expense of thermic energy; on the contrary, heat as well as work is increased with the heavier weight. The second fact becomes quite obvious if thus regarded:—like caoutchouc, muscle is warmed by passive elongation, and if, of two identical and equally weighted muscles, one pulls up and lets down



its load (*i.e.* no work done), while the other winds up its load from a lower to a higher level (*i.e.* work done), the first muscle, alternately contracting and stretched, must be warmed more than the second muscle, that only contracts, and is not stretched.

As we have seen above, the muscles, even at rest, are constantly slightly warmer than arterial blood, and the difference is increased during contraction. Some idea of the amount of heat evolved by muscular contraction may be arrived at by comparing the temperature of the venous blood coming from tetanised muscles with that of arterial blood. If, for instance, through a given group of muscles 100 c.c. of blood have passed per minute, and have been raised to a temperature  $0.2^{\circ}$  above that of arterial blood, it follows that heat has been produced at the rate of 20 calories per minute in the muscles in question. Such an estimate does not, however, give the total amount of heat evolved; to it must be added the value of any rise of temperature of the muscle itself. If, for instance, the muscle in action weighs 50 grms. and its temperature is raised  $0.1^{\circ}$ , this value will be about 5 calories—to be added to the 20 calories as above estimated. And even then we have left out of account the heat dissipated by radiation from the muscle.

Direct estimations of the amount of heat evolved have been obtained on excised frog's muscles, and on mammalian muscle after the circulation has been arrested. Fick estimated that frog's muscle evolves 1 to 3 milli-calories per 1 gramme in a single contraction. Ludwig gives for dog's bloodless muscle a heat production of 1 milli-calorie per gramme per contraction. It is probable that these values are below the value of normal heat-production of muscle in the body, more particularly in the case of mammalian muscle, which rapidly runs down after arrest of the circulation. Chauveau's observations performed under more normal conditions, upon the levator labii superioris of the horse, yielded a much higher value, viz., 300 milli-calories per gramme per minute during voluntary contraction.

Briefly recapitulated, the following five propositions express the principal results of heat-experiments on isolated muscle:—  
1. Muscular contraction produces heat (Helmholtz). 2. Interrupted contraction produces more heat than does continuous contraction (Fick). 3. Less heat is produced by a muscle that does 'work' than by a muscle that does no work (Fick). 4. The greater the tension against which muscle contracts, the greater is the heat evolved (Heidenhain). 5. In fatigue the

thermogenic power declines more rapidly than the height of lift (Heidenhain).

The production of heat by muscular exercise is a matter of everyday experience, and the increased temperature of a single limb (*e.g.* the arm or the forearm) in consequence of muscular contraction is easily demonstrated on man by means of an ordinary thermometer with tenths of a degree legible; the temperature may rise as much as  $1^{\circ}$  or even  $2^{\circ}$  in consequence of strong muscular contractions repeated for two or three minutes. But this rise is not a pure muscular effect; it is mainly due to a more copious blood-supply, as may be proved by repeating the observation during arrested circulation (by a bandage); the rise is then very much smaller than that just obtained with unobstructed circulation. This result is in apparent contradiction to the statements just made with reference to the higher temperature of muscle as compared with arterial blood. But these statements refer to 'deep' temperature of well-protected muscle, whereas on man the temperature taken is of superficial tissues (skin and subjacent muscle) that are cooled below blood-temperature, and therefore warmed by more copious blood-supply. The warming thus effected varies with the magnitude and duration of muscular effort; in an experiment on the forearm the rise of temperature was, *e.g.*,  $0.75^{\circ}$  after 15 contractions of 10 kgs.,  $1.1^{\circ}$  after 30 contractions of 10 kgs.,  $1.4^{\circ}$  after 15 contractions of 20 kgs. (w.)

**The three lever systems.**—The mechanical disposition of the muscles, tendons, and bones of the animal body presents examples of the three fundamental lever systems. As instances of the first system, in which the *fulcrum* is situated between the *power* and the *weight*, may be mentioned the action of the triceps brachii in the extension of the forearm, that of the hamstring muscles in raising the body from a forward stoop, that of the calf muscles in the movement of pointing the toe; as an instance of the second system, in which the weight is placed between the fulcrum and the power, may be mentioned the action of the calf muscles in raising the body on tiptoe; as instances of the third class, in

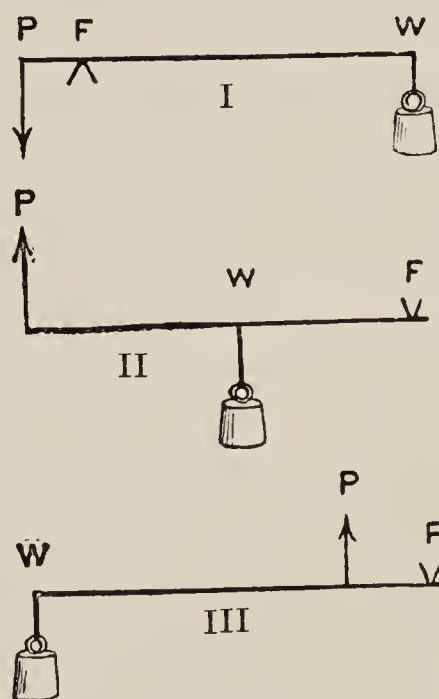


FIG. 153.



which the power is applied to a lever between the fulcrum and the weight, may be mentioned the actions of the biceps upon the forearm, of the hamstrings upon the leg, of the temporal and masseter muscles upon the lower jaw. Levers of the second system afford increase of *power* at the expense of quickness and range of movement, the power-arm of the lever being longer than the weight-arm, as is illustrated in the mechanism of a nutcracker; levers of the third system afford increased *quickness* and range of movement at the expense of power, the weight-arm of the lever being longer than the power-arm, as is illustrated in the use of a fishing-rod or a golf-club. Levers of the first kind may afford either of these characters at the expense of the other, according as the distance F P is greater or smaller than the distance F W; generally, however, the mechanical conditions in the body are such that this system affords quickness levers rather than power levers. The fuller and more detailed consideration of the principal lever movements of the body in various circumstances belongs to applied anatomy rather than to physiology.

**Co-operative antagonism.**—It is a fundamental principle common to the neuro-muscular actions of the body, that the normal voluntary movement of any given group of muscles is usually associated with *contraction* of the antagonistic muscles. This physiological principle of ‘co-operative antagonism’ may be easily realised by simple observations, is illustrated in familiar pathological occurrences, and offers analogies outside the domain of medicine, beyond which we shall, however, not pursue it. To take a simple and easily observed case, we may refer to the flexors and extensors of the fingers; if the fingers of one hand be flexed and the forearm felt by the other hand, it will be noticeable that the contraction of the flexor muscles is accompanied, not by relaxation, but by *contraction* of the extensors; and when the fist is clenched, it will be still more obvious that contraction of the flexors is accompanied by *contraction* of the extensors.

That to the effective action of a muscular group the effective action of opposed muscles is necessary, is well illustrated in *lead-palsy*; in a typical case of this affection the hand drops at the wrist in consequence of paralysis of the extensor muscles; the flexors are not paralysed, but movements of flexion cannot be carried out. Conversely, cases are met with of muscular paralysis limited to the flexor group of forearm muscles, when extension, as well as flexion, become impossible—*i.e.* to the due execution of a measured movement either of flexion or of extension, the

integrity of the antagonists as well as of the direct motors is necessary; or otherwise, antagonistic muscles co-operate in the execution of movements, now one, now the other group exercising the major or predominant power. This is the rule, but there are exceptions to it; in very rapid movements, and in movements against an insuperable external resistance, the contraction of groups of muscles is not accompanied by any contraction of the antagonists, which may even relax. It is a subordinate although obvious feature of any extensive movement, that in addition to the *active* participation of antagonists, these must also undergo a *passive* elongation, calling into greater play their physical property of elasticity, and thus contributing to the steadying effect that is an essential condition of all delicate adjustments.

**Curare** is the drug most frequently spoken of in association with muscle and nerve. A curarised frog lies motionless and paralysed, yet its muscles are excitable, and so are its nerves; a curarised cat, or dog, or rabbit, is equally paralysed, and dies by arrest of respiratory movements unless artificial respiration is employed.

The fundamental experiment demonstrative of the action of curare is as follows:—a drop of a 1 per cent. solution is injected under the skin of a frog, and when the animal is found to be completely paralysed, it is pithed and the sciatic nerve is exposed, or a nerve-muscle preparation is put up as shown in fig. 138. Excitation of the nerve produces no effect, while the muscles are excitable as before. By other experiments it may be shown that the nerve retains excitability, *e.g.* the negative variation persists as before. Seeing that the muscle is excitable, that the nerve is excitable, but that excitation of the nerve fails to put the muscle in motion, it is inferred that curare acts upon the junction between nerve and muscle, *i.e.* that it causes a block at the motor end-plates.

It may be shown by a variation of the experiment that curare has practically no action upon sensory fibres or cells. A frog is curarised as above, but one limb is protected by a tight ligature leaving out the nerve: when the paralysis is complete it will be found that cutaneous stimulation of any curarised part causes reflex movements of the protected, and therefore uncurarised limb.

**Ciliary movement.**—Cilia—the sole means of locomotion in some of the lower animalcules (*e.g.* ciliated infusoria)—are in the higher animals confined to certain surfaces and ducts where



they apparently assist in the extrusion of solid particles. They possess a spontaneous contractility independent of any direct influence of the nervous system, subject only to the vitality of the cells with which they are connected, and therefore to temporary suppression by carbon dioxide and other anæsthetics.

In man their chief situations are the respiratory passages, the tympanum and eustachian tube, the uterus and the fallopian tubes, the testicular ducts, the central canal of the spinal cord and the ventricles of the brain. In these last situations their presence is enigmatical unless they be regarded as signifying that an ancestral gut has found its present representative in the cerebro-spinal axis. As regards the other situations it may be observed that the sharp effective stroke of the cilia is expulsive, and that the collective force exerted by a field of cilia is surprisingly great. One of the most convenient objects for the study of ciliary movement is supplied by the frog's gullet; a shred of its mucous membrane laid cilia uppermost on a glass slide may be seen to clear itself of quite 'large' blocks of foreign matter (microscopically regarded), or turned cilia downwards the whole shred may crawl along the slide; Engelmann has actually made a light wheel to be driven by ciliary movement and to raise a weight, thereby affording the possibility of approximately estimating work done.

## CHAPTER X

## THE PERIPHERAL NERVOUS SYSTEM. NERVE

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- 347 Classification and terminology—Minute structure.
- 353 Chemical composition.
- 354 Section and excitation—‘Law of specific nervous energies’—‘Law of isolated conduction.’
- 356 **Consequences of nerve-section:** (1) Paralysis—(2) Loss of excitability—(3) Degeneration—(4) Regeneration—(5) Restoration of function and of excitability.
- 356 **Degeneration:** of mixed nerve—of anterior root—of posterior root—‘Trophic centres’—Immediate restoration (?)—Recurrent sensory fibres.
- 359 \* **Trophic nerves** (?)
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- 364 \* **The constant current:** Make and break contractions—Pflüger’s law of contractions—Ritter’s tetanus—Formula of contractions on man—Reaction of degeneration.
- 369 \* **Electrotonus:** Electrotonic currents—Electrotonic alterations of excitability—The paradoxical contraction—Experiments on man—Experiments on sensory nerves.
- 377 **Velocity of nervous impulses:** In motor nerves of man—of frog—In sensory nerves (?)—In sympathetic nerves (?)—Velocity of negative variation.
- 379 Conduction in both directions—Paul Bert’s experiment—Du Bois-Reymond’s experiment—Kühne’s experiment.
- 380 Inequalities of excitability—At different points—Of different fibres. Avalanche theory—The Ritter-Valli ‘law’—In motor nerves—in sensory nerves.
- 382 **Fatigue** of nerve—of muscle—of the motor end-plate—Inhibitory stimulation.

*Classification and terminology.*—Nerves and nerve-fibres are classified from several points of view—(a) according to their obvious source and course, as *cerebro-spinal* and *sympathetic*; (b) according to their microscopical structure, as *medullated* and *non-medullated*; (c) according to their embryonic origin and distribution, as *somatic* and *splanchnic*; (d) according to their function, as *efferent* and *afferent*. And we shall learn shortly that these divisions are subdivided into several varieties according to the kinds of impulses conveyed. As regards the correspondence of the various kinds denoted by these terms, it is to be remarked,



(1) that cerebro-spinal nerves are in major part composed of medullated fibres, in minor part of non-medullated fibres, while sympathetic nerves are in major part composed of non-medullated fibres, in minor part of medullated fibres; (2) that both medullated and non-medullated fibres may be either afferent or efferent, there being here no correspondence between difference of structure and difference of function. The interchange of fibres between the two classes of nerves is by the *rami communicantes*, through which the cerebro-spinal nerves receive grey fibres from the sympathetic system and give to that system white fibres. But, as we have already seen, no hard-and-fast line of distinction is to be drawn between the two systems. Sympathetic nerve-fibres have their centres in the spinal axis, and not in their own ganglia.

A certain amount of correspondence is presumed to exist between the size of fibre and the nature of its function. As a rule, the largest nerve-fibres (15 to 20  $\mu$ ) are those distributed as motor fibres of skeletal muscle; fibres in the posterior nerve-roots and in the posterior columns of the cord are on the average of inferior calibre to fibres in the anterior roots and in the pyramidal tracts. Another difference has already been alluded to under the physiological anatomy of cardiac and vasomotor nerves, where we saw that these are characterised by their very small diameter (2 to 4  $\mu$ ), and stated that in their course along the peripheral nerves vaso-inhibitory fibres are medullated, while vaso-augmentor fibres are non-medullated.

Medullated or white nerve-fibres owe their name to the medulla or myelin, which is a fatty sheath surrounding the cylinder-axis. From within outwards a fibre of this kind consists of—(1) axis-cylinder; (2) medullary sheath or white substance of Schwann; (3) primitive sheath or neurilemma.

The axis-cylinder is the essential or conducting part of the fibre; it is to be regarded as a continuous filament of protoplasm formed (according to Engelmann) by long segments very perfectly joined end to end. The medullary sheath is a string of hollow cylinders of a fatty nature surrounding the axis-cylinder and joined end to end by cement substance at 'nodes' (Ranvier); this is an indication that each internode constitutes a single cell, a view which is borne out by the fact that each such internode possesses one, and only one nucleus. The neurilemma is to the nerve-fibre what the sarcolemma is to the muscle-fibre, viz. a delicate membranous sheath enclosing its substance. Outside this a second membranous sheath is commonly visible, enclosing solitary

nerve-fibres ; this is Henle's sheath, and is not an integral part of the fibre, but a prolongation derived from the connective tissue (perineurium), by which nerve-fibres are held together in bundles.

A 'non-medullated,' 'pale,' or 'grey' fibre is nerve reduced to its simplest expression—a naked axis-cylinder. It is to be regarded as being less highly developed than medullated nerve, of which it is the precursor in the embryo : in adult life it exists in greatest abundance in visceral nerves, in association with non-striped muscle, and histologically, in addition to its simpler structure, it is characterised by the presence of numerous nuclei.

*Motor and efferent*—*sensory and afferent* are not completely synonymous pairs of terms. All motor nerves are efferent, but all efferent nerves are not motor ; some are anti-motor or 'inhibitory' of motion. Sensory nerves are afferent, but all afferent nerves are not necessarily 'sensory' in the exact

acceptation of the term, 'sensory' being strictly applicable, not to all centripetal impressions, *but only to such as excite consciousness*. Many, if not all, afferent channels may, however, on some occasions convey impulses that reach consciousness, while habitually conveying impulses of which we remain unconscious. Normally, we are not conscious of the beating of the heart, nor of digestive actions ; in disease we may become painfully conscious of them. On the other hand, all, or at least most, sensory channels may on some occasions convey impulses of which we remain unconscious, though habitually we perceive them. A man may fail to perceive that he has received a serious injury if his perceptive centre is preoccupied by a sufficiently powerful idea or sensation. As regards the distribution of these several kinds of nerves, cerebro-spinal nerves are distributed for the most part to the skin and to voluntary muscles, sympathetic nerves for the most part to viscera and blood-vessels ; skin-nerves are mostly sensory in function, muscle-



FIG. 154.

Transverse and longitudinal sections of a medullated nerve-fibre.



FIG. 155.

Longitudinal optical section of a group of non-medullated fibres.



nerves are mostly motor in function. The skin, however, contains glands and a small proportion of unstriated muscle, which receive motor nerve-fibres derived from the sympathetic, causing contraction or dilatation of its vessels, erection of its hairs, secretion of its glands, shrinking of its pigment-cells; and voluntary muscles, or more precisely their tendons and sheaths, possess also a small proportion of sensory nerves, through which the sensorium is kept informed of their state. Motor nerve-fibres are also distributed to internal organs: the impulses that they convey are, however, independent of the will; by voluntary effort we cannot directly move the involuntary muscle of such organs.

**Nerve-fibres** may be classified as follows:—

AFFERENT.	{	olfactory . . . . .	{	cutaneous sensory			
		optic . . . . .	{	muscular sensory			
		gustatory . . . . .	{	visceral sensory			
		auditory . . . . .	{	? thermic sensory			
		tactile . . . . .	{	? pathic			
EFFERENT	{	MOTOR	musculo-motor . . . . .	to skeletal muscle			
			vasomotor . . . . .	{	to cardiac muscle (accelerators)		
				{	to arterial muscle (constrictors)		
			pilomotor . . . . .	to muscle of hair			
			broncho-motor . . . . .	to bronchial muscle			
			viscero-motor . . . . .	to intestinal muscle			
			secreto-motor . . . . .	to gland cells			
			? 'thermogenic' and 'trophic'	{	to all tissues?		
					INHIBITORY	vaso-inhibitory	{
		{				to arterial muscle (dilating)	
		viscero-inhibitory	to intestinal muscle				

The most prominent manifestation of function subserved by efferent nerves is voluntary muscular movement; there is as yet no proof of direct voluntary inhibition at the periphery; voluntary inhibition of a peripheral action occurs in the centre, whence the action would have been excited in the absence of inhibitory control. Of involuntary motor paths we have proof in the action of nerves upon arterial muscle and upon secretory glands; both these are entirely exempt from direct voluntary influence; all that we can voluntarily effect is to apply a peripheral stimulus that will influence them by a reflex mechanism. *Vaso-constriction* is a motor effect; *vaso-dilatation* is of an inhibitory nature;

we do not know, however, whether the seat of inhibition is in the muscular element itself, or in some hypothetical peripheral centre contained in it. It is analogous with the better-known case of *cardiac inhibition* by the *vagus*, but in this case also we do not know whether the motor action is interfered with in the motor element itself, or in an intermediate nerve-element—the ganglion cell. Closely analogous with vasomotor and vaso-inhibitory nerves, but still less completely understood, and, indeed, less well certified by experiment, are the effects upon *intestinal movements* of impulses passing to them through efferent channels. Intestinal movements may through such channels be *excited* or *arrested*; usually, the result of *vagus* excitation is increased movement, of *splanchnic* excitation diminished movement; but such results are not infallibly obtained, being dependent upon a variety of circumstances (see p. 168).

*Secreto-motor* fibres are subdivided into two varieties: 1, such as accelerate the discharge of water; 2, such as accelerate the discharge of the protoplasm-product, which is to form the essential constituent of the secretion. These latter secreto-motor fibres are distinguished from the former as ‘trophic,’ a term which is not free from objection, seeing that such fibres have not been proved to excite cell-nutrition, though they certainly effect the converse, viz. cell-disintegration (see p. 182).

*Does muscle possess inhibitory nerves?* A muscle can at will be relaxed as well as contracted, and it may and has been asked whether we must attribute the relaxation to an action of arrest in the brain, or to an inhibitory effect in the muscle by special anti-motor nerve-fibres. We shall more conveniently discuss this second alternative at a later stage (p. 384), in considering the relaxant effects upon muscle of experimental stimulation of its nerve. Meanwhile, as regards voluntary arrest of action, it will be sufficient to state that there is as yet no proof of direct voluntary inhibition at the periphery, but that voluntary inhibition of a peripheral action occurs at the centre (cortical or bulbar) whence the action would have been excited in the absence of inhibitory interference.

**Afferent nerves** are naturally classified in accordance with their obvious functions as *olfactory*, *optic*, *acoustic*, *tactile*, and *common sensory*. There are less obvious and less satisfactory reasons for admitting a further classification of nerve-fibres into *thermic* and *pathic*, for the proof that impressions of *heat*, of *cold*, and of *pain* are specific and served by separate fibres, distinct from common



sensory fibres, is not completely clear and unassailable. The term 'sensory' is in common use, and cannot be advantageously ignored; it is well, therefore, to recognise clearly that the same nerve or the same nerve-fibre may at one time be truly sensory, inasmuch as it may be the channel of a peripheral stimulus that excites consciousness, while at another time it may be only the channel of an impulse that effects a reflex act in the absence of consciousness. There is no reason for believing that reflex acts are effected through nerve-fibres, afferent or efferent, other than those which convey sensory and voluntary impulses; the same afferent and efferent fibres convey reflex involuntary impulses (which may or may not be perceived) to and from the spinal cord, as well as true sensificatory impulses and willed motor impulses to and from the brain, *via* the bulb and cord. Further, it may be repeated, that whereas some nerves usually carry an impulse to its supreme end as a sensation, while others usually carry impulses short of this, so that 'sensory' is the term applied to the former, 'afferent' to the latter—yet the distinction thus implied is not real, for any such habitually 'sensory' nerve may on occasions be merely afferent, and, *vice versa*, any such habitually 'afferent' nerve may on occasions be sensory.

It will also have been noticed in the tabular summary that whereas efferent nerves are either motor or inhibitory, we have not explicitly recognised the same distinction in the case of afferent nerves. Any afferent nerve is, in fact, both motor and inhibitory as regards its central effects; its excitation raises excitability at one part, while it simultaneously depresses it at other parts of the central nervous system. This double action will be better appreciated when the phenomena of hypnosis have been considered. Moreover, the same afferent nerve acting upon the same excitable central part at different times may now raise excitation of a depressed part, now depress excitation of the excited part. In Brown-Séquard's language, it may have either a dynamogenic or an inhibitory effect according to circumstances.

*Does muscle possess sensory nerves?* In giving an answer to this question we must distinguish between the muscle itself and its connective tissue. Nerves are distributed in tendon, and their terminations are fully described. Tendons are not contractile, their nerves cannot be motor, and must therefore be sensory, or at least afferent. Moreover it is known that the extension of muscle is felt, and we have reason to believe that

nerves from tendon are the centripetal channels of the effect. But with regard to the muscle itself there is no absolute anatomical proof of the existence of sensory fibres; on the frog, after section of the posterior roots through which all afferent channels pass, all such fibres degenerate beyond the point of section, and no degenerated fibres have been found in the muscle itself. In sum it is proved that tendons possess afferent fibres, it is not proved that muscles possess afferent fibres; so that the proper answer to the paragraph heading is No, as regards the muscular substance Yes, as regards the connective tissue of muscle.

The classification into 'splanchnic' and 'somatic'<sup>1</sup> is derived from the corresponding embryological terms denoting the two layers of the mesoblast that respectively become viscus and body-wall. The terminology, as applied to nerves, does not imply that they are developed from the splanchnic and somatic layers of the mesoblast (all nerves, cerebro-spinal and sympathetic alike, being outgrowths from a central axis which is epiblastic), nor even that they supply tissues which are derived from these two layers respectively, seeing that a 'somatic' tissue, *e.g.* the skin, receives 'splanchnic' (vasomotor and secreto-motor) as well as somatic (sensory) nerves.

The origin of the terms as used by Gaskell is as follows:—*a.* The vascular, intestinal and glandular nerves were classed under the designations 'visceral' or 'splanchnic.' *b.* Embryological descriptions were taken to justify the recognition of Bell's respiratory system of nerves (facial, phrenic, intercostals, spinal accessory) as 'splanchnic,' and as being derived from a separate 'lateral root,' supplying the lateral or ventral plates of the mesoblast (which form part of the somatopleure, as well as of the splanchnopleure), as distinguished from the axial or dorsal portion forming the mesoblastic somites. *c.* This 'lateral root' hypothesis was extended to the whole cord and brain, and made to include all vascular, visceral, and glandular nerves. Thus Gaskell's 'somatic nerves supply parts derived from the epiblast and from the mesoblastic somites,' his 'splanchnic nerves supply parts derived from the hypoblast and from the rest of the mesoblast.' The latter comprise the vascular, visceral, glandular, and respiratory nerves.

The chemistry of nerve is very imperfect, and from a physiological standpoint very insignificant—no differences having ever been detected between living and dead, or between rested and exhausted, nerve. The only point which appears to be established is that normal nerve-matter is faintly alkaline, while immediately after death it is found to be acid; it is probable,

<sup>1</sup> σπλάγχνον, viscus; σῶμα, body.



though not proved, that acidification also takes place during life in the cerebral grey matter, as a consequence of activity.

Our scanty knowledge of the subject is based upon analyses of the white matter of the brain, and upon micro-chemical reactions. From the first source it has been ascertained that 'nerve-fat' is of a peculiar composition, and that it contains cholesterin, lecithin, cerebrin, and protagon.

*Cholesterin*, with which we have become acquainted as a constituent of bile, forms at least one half of this so-called fat, and the coincidence recalls to mind the everyday experience that liver action has influence upon brain action. No definite relationship has, however, been determined, and the supposition that the blood can be overcharged with cholesterin so as to give rise to a definite group of symptoms (cholesteræmia) has not been verified. *Lecithin* is the essential constituent of the so-called 'myelin' or white substance of medullated nerve, which is fixed and blackened by osmic acid. *Cerebrin* is the name given to the indefinite phosphor-free body or group of bodies obtainable from brain-matter, probably by decomposition of 'protagon.' The crystalline phosphorised body discovered by Liebreich and termed *protagon*, but considered by Hoppe-Seyler as a mixture of cerebrin and lecithin, is probably a compound of these bodies, *i.e.* a true proximate principle (Gamble). Kühne has described, under the name of *neurokeratin*, the reticular formation which he found to pervade the medulla of nerves, and to resist the action of the digestive ferments. The *neuroglia* of nerve-centres is of a similar character.

Analysis of the white matter gives on the average :—

Water . . . . .	70 per 100
Solids { Cholesterin . . . . .	15.0
{ Proteids . . . . .	7.5
{ Lecithin . . . . .	3
{ Cerebrin . . . . .	3
{ etc. . . . .	1.5
	30 „

The **function** of a nerve is ascertained by the observation of the alterations consequent upon (1) its *section*, (2) its *excitation*, such excitation being applied to the central and to the peripheral cut ends. The immediate consequence of nerve-section is 'paralysis'—in the case of motor nerves, inability to move the muscles served by the cut nerve—in the case of sensory nerves, inability to perceive sensations from the sensory district served by such nerves. All nerves may convey *occasional* impulses; some nerves are *constantly* engaged in the conduction of centri-

petal or centrifugal impulses, and the actions thus effected are spoken of as *tonic*. An instance of such tonic action in the case of an efferent nerve is furnished by the cardiac fibres of the pneumogastric, in the case of an afferent nerve by the pulmonary fibres of the same nerve. If nerves that convey such tonic influence be divided, the effect is to cut it short; thus, after section of the pneumogastrics the heart's beat is accelerated, the respiratory movement is slowed; these results, taken in conjunction with the confirmation obtained by excitation of the central and peripheral ends of the nerve, justify the conclusion that it conveys tonic influence to the heart, which restrains its action, tonic influence from the lung, which accelerates respiratory action at its seat of government—the spinal bulb.

The proof of the function of a nerve is completed by excitation, which is to be successively applied to the two ends of the divided nerve. In the case of most nervous impulses, which are not tonic but only occasional, excitation after section, and not section alone, furnishes the proof of function. Excitation of the peripheral end of a nerve, followed by an effect at the periphery, proves that the nerve contains efferent fibres, and the kind of effect at the periphery, whether motor, or inhibitory, or vasomotor, shows what kind of efferent fibres the nerve contains. The absence of effect at the periphery proves the absence of efferent fibres; if such absence of effect is witnessed on a freshly-divided nerve, efferent fibres are really absent from the normal nerve; if it is witnessed on a diseased nerve, or on a nerve divided some time previously, efferent fibres are *functionally* absent from the nerve, which has been rendered abnormal (degenerated) by previous interference or by disease. Excitation of the central end of a nerve, followed by movements expressive of sensation, or by vasomotor reactions, proves that the nerve contains afferent fibres.

It is known from observations made upon man that excitation of the central cut end of a sensory nerve excites in consciousness the kind of sensation which that nerve habitually subserves. This is what is meant by the term '*law of specific nervous energies*.' A second important fact in nervous action, viz. that nerve-fibres constitute uninterrupted channels between central and peripheral organs, isolated from adjacent fibres, is spoken of as the '*law of isolated conduction*.' Both these 'laws' are implied in the familiar phrase, 'each nerve-fibre minds its own proper business.' In general it is excitation of the central cut end of a nerve that excites sensation, excitation of the peripheral end



being unfelt. But the rule is not absolute ; the peripheral ends of many mixed nerves contain sensory fibres along which the impulses run towards the periphery before returning to their central destination. Such afferent fibres are known as *recurrent sensory fibres* ; they are demonstrable in the peripheral end of the divided anterior root of spinal nerves and in the peripheral ends of mixed nerves (pp. 359, 486).

**The consequences of nerve-section.**—Division of a nerve has for its consequences in order of time—(1) *paralysis* of motion or of sensation, or of both, according as the nerve cut is motor or sensory or mixed ; this paralysis is immediate and local ; (2) *loss of excitability* of the nerve, coming on gradually and becoming complete within a few days, direct muscular excitability persisting for an indefinite period, especially to the galvanic current ; (3) *degeneration* of the peripheral end of the nerve, also a gradual process, visible within a day or two, well marked at the end of three or four days, complete in about ten days ; (4) *regeneration* of the previously fully-degenerated peripheral end of the nerve, a still more gradual process, commencing indefinitely, but clearly

visible about a month after the lesion has been produced, requiring at least six months to complete itself, and resulting in (5) *restored motility, sensibility, and excitability*. These statements refer to the results of experiments upon warm-blooded animals ; upon cold-blooded animals the last four stages are much more slowly consummated ; upon young warm-blooded animals, on the contrary, the processes are accomplished more rapidly.

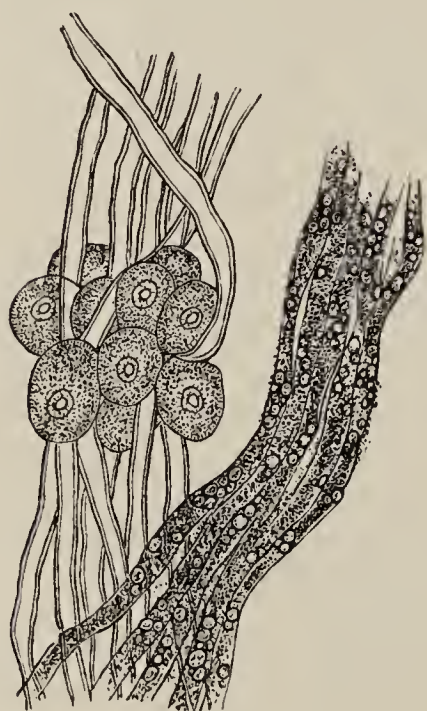


FIG. 156.

Groups of fibres from the anterior and posterior roots several days after section of both roots close to the cord ; the anterior fibres are degenerated, the posterior are normal. (From an unpublished drawing by A. Waller, 1852.)

**Trophic action of nerve-cells upon nerve-fibres.**—*Wallerian degeneration*.

The degeneration of nerve after section is a coarse and easily recognised change ; hence the practical value of the method in the investigation of the distribution of nerves. A mixed nerve degenerates down to its ultimate distribution in muscle or in skin ; the process is simultaneous throughout the length of the nerve, and does not gradually descend towards the periphery, there being no difference recognisable between fibres in a nerve-

trunk near and far from the section; according to Ranvier, the earliest signs of change are detected in the intramuscular nerve-ends. The rapidity with which the process takes place varies with the activity of tissue-life—*e.g.* it is more rapid in a young than in a full-grown animal, in a cold-blooded animal kept at a high temperature than in one kept at a low temperature, in a healthy than in an enfeebled animal. The histological features are significant of excessive rather than of deficient metabolism, the nerve-cell being, so to speak, the nurse of the nerve-fibre, so that if the fibre is separated from the cell, it runs riot and destroys itself. The entire process is naturally divisible into two stages—(1) the period preceding the loss of excitability, *i.e.* under ordinary circumstances up to about the third or fourth day; (2) a succeeding period of three or four weeks. During the first period the signs are those of increased activity—hypertrophy of protoplasm, hypertrophy and multiplication of nuclei, causing interruptions of the medullary sheath and of the conducting medulla itself. During the second period absorption of the broken-up fibres takes place, leading to their complete disappearance, and leaving nothing but a strand of connective tissue, which probably includes the actual primitive sheaths of the vanished fibres. Regeneration—the first signs of which are visible during the actual process of degeneration—occurs from the central end, and pushes from it to the periphery, being a repetition of the original process of nerve-development. The axis-cylinders of the central end hypertrophy, and if examined two or three months after section, are found to have given off fine pale fibres from their club-shaped ends; these grow in length and breadth, insinuate themselves along the old track, develop a medullary sheath, and become new nerve-fibres; 1 mm. per day (according to Vanlair's observations on dog's vagus and sciatic) represents an average rate of regrowth. It remains to be stated that this process of degeneration and regeneration of nerve is to some extent a normal physiological phenomenon: the same nerve-fibres do not persist indefinitely during life, but some are dying while others are growing up; in a normal nerve, on careful examination, a few degenerating nerve-fibres are discoverable, as

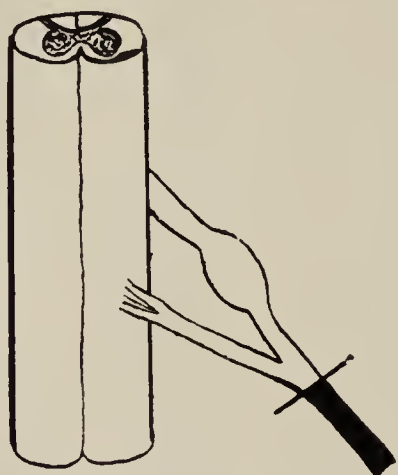


FIG. 157.—REGENERATING NERVE-FIBRES; MONTHS AFTER SECTION. (Ranvier.)

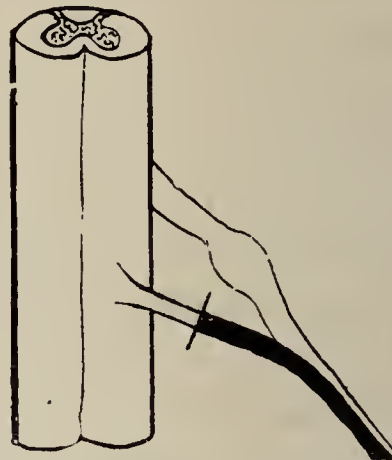


well as numerous fine fibres, non-medullated as well as medullated, which not improbably include young fibres (S. Mayer).

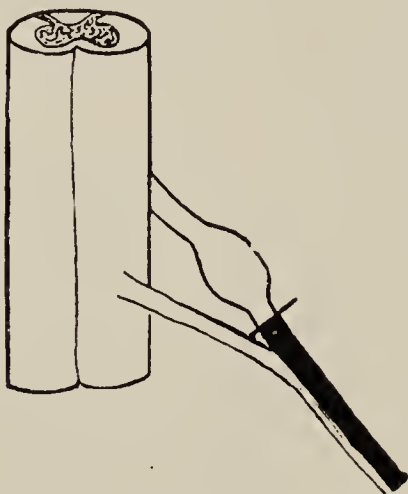
*Primary union* of nerve, with immediate restoration of conductivity, has never been experimentally realised, and the cases on record of return of sensibility or of motility a day or two after accidental division and surgical reunion of nerves, cannot be received as evidence to counterbalance the undoubted fact



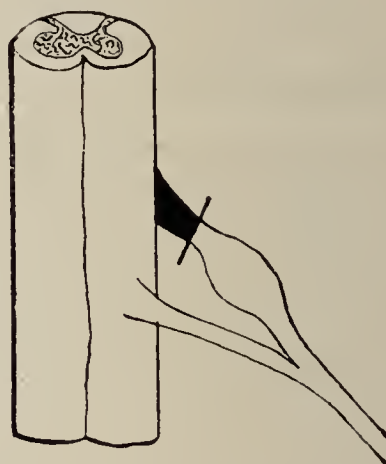
Degeneration of efferent and of afferent fibres below a section of entire nerve.



Degeneration of efferent fibres below a section of anterior root.



Degeneration of afferent fibres below a section of posterior root beyond the ganglion.



Degeneration of afferent fibres above a section of posterior root above the ganglion.

FIG. 158.—DIAGRAMS TO ILLUSTRATE WALLERIAN DEGENERATION OF NERVE-ROOTS.

that any section, however carefully the cut ends are brought together, entails degeneration in the peripheral end, and loss of function, which require *months* to be recovered from. It is more probable in such cases that the nerve has been temporarily compressed, or that recurrent sensory fibres have recovered excitability after a temporary depression by 'shock,' or that the clinical examination has been illusory.

Attempts to join motor with sensory nerves—*e.g.* the central end of the hypoglossal with the peripheral end of the lingual—and to show subsequently that a former sensory nerve produces motion, and a former motor nerve sensation, have furnished no

evidence of any value. But the attempt in the hands of Phillipeaux and Vulpian led to the discovery of an extremely paradoxical effect, produced by excitation of the peripheral end of the lingual nerve after degeneration of the hypoglossal, viz. a very sluggish contraction of the lingual muscle. This contraction is due to the chorda tympani nerve, and is accompanied by vaso-dilatation, but its mechanism is quite incomprehensible—the chorda has no anatomical connection with the lingual muscle, and produces no effect with intact hypoglossal nerve; Heidenhain has termed it a pseudo-motor effect.

It is the entire peripheral end of a mixed nerve that degenerates, *i.e.* motor and sensory nerves degenerate irrespectively of any direction of function in them. If the anterior root of a spinal nerve be divided, the degeneration is of the peripheral end. If the posterior root of a spinal nerve be divided *below* the ganglion, the degeneration is likewise of the *peripheral* end. But if the posterior root be divided *above* the ganglion, the degeneration is of the central end. From these facts it is concluded that the normal nutrition of motor nerve-fibres is dependent upon the grey matter of the spinal cord, while that of sensory nerve-fibres is dependent upon the ganglion of the posterior root. The grey matter of the cord (*i.e.* the cells of the anterior cornua) and the ganglia of the posterior roots are therefore spoken of as the ‘*trophic*’ centres of the fibres of the anterior and of the posterior roots respectively.

These are the fundamental facts; two supplementary points should, however, be mentioned. (1) After the anterior root has been divided, its peripheral end degenerates with the exception of a few fibres which remain normal, and the central end contains a few degenerated among the large majority of normal fibres: after the posterior root has been divided below the ganglion, a few degenerated fibres are discoverable in the anterior roots. These few exceptional fibres are the recurrent sensory fibres alluded to below (p. 486); they have their trophic centre in the ganglion of the posterior root. (2) According to Max Joseph, there are also in the posterior root a few fibres that escape the general degeneration on their separation from the ganglion, and that presumably have their trophic centre in the cord or at the periphery, simply traversing the ganglion without connection with its cells.

**Trophic nerves.**—There is abundant evidence to show that nerves can influence the nutrition of tissues, but whether they do



so by a direct trophic action, or indirectly by causing vascular modifications, or otherwise, is the question we have now to examine. We have positive evidence that in the nervous system itself nerve-cells exercise a trophic influence upon nerve-fibres, but this definite knowledge nowise includes or answers the question we are now putting, and which we may formulate as follows :—Are nerves in general, or in any one indisputable case, the channels of an influence directly modifying the nutrition of peripheral tissues, independently of vascular or other changes? Is there any proof of the existence of ‘trophic’ as distinguished from ‘motor,’ ‘secretory,’ or ‘inhibitory’ nerves?

The cases in point that naturally come first to mind are—(1) the effects of division of the fifth nerve; (2) the effects of division of the cervical sympathetic, and of other nerves in young, rapidly growing animals; (3) clinical incidents, especially the appearance of ‘bed-sores’ and of ‘glossy skin’ on paralysed parts, and of herpes along the course of cutaneous distribution of posterior nerve-roots.

The last-named items, however interesting in themselves, are in reality not admissible in either an affirmative or a contradictory sense as regards the answer to the questions stated above. No doubt the nutrition of paralysed parts is altered, and bed-sores develop with surprising rapidity; equally surely, in the case of herpes, a trophic disturbance coincides with the distribution of nerve-roots; we have no proof that the malnutrition is a direct trophic effect, to the exclusion of the unnoticed rough usage or prolonged pressure to which anæsthetic parts are liable, or—in the case of herpes—to the exclusion of vasomotor changes; but it is a remarkable fact, almost amounting to proof of a real trophic disturbance beyond what might be attributed to coarse injury, that a paraplegic dog resting on the ventral surface develops a dorsal sore analogous in situation with the ‘bed-sore’ that is apt to occur in a paraplegic man.

Turning next to experimental data, we have also to dismiss as imperfect evidence the greater growth of a young rabbit’s ear after section of the cervical sympathetic, the greater growth of the maxillary bones after section of the facial nerve, and the wasting of an inferior extremity after section of the sciatic nerve; the first two instances are obviously attributable to increased vascularity of the parts, the third is the extremely gradual effect of paralytic disuse of the limb, or an acute effect of which the intimate mechanism is most obscure; an acute muscular atrophy of such character, with ‘reaction of degeneration’ (p. 369), is

perhaps the nearest approach we possess to evidence of the existence of trophic nerves, but it is not 'clean' evidence. We cannot say whether the effects are paralytic or irritative, nor whether they are direct or indirect or vascular, and there are discrepancies between the clinical and the experimental phenomena upon which we cannot enter here, but which must be cleared away before the evidence can be admitted as unequivocal.

The next case we have to consider is that of the fifth nerve, to which, more than to any other nerve, appeal has been made by experiments in this connection. Shortly, the facts are these: after section of the fifth nerve the eye becomes inflamed, the cornea ulcerates, the entire eyeball suppurates and is lost. To the conclusion that these are an instance of trophic disturbance, objection has been raised to the following effect:—After section of the nerve, the conjunctiva and the cornea lose sensibility, foreign bodies lodge themselves unnoticed on the eyeball, and give rise to inflammation. In support of this interpretation it has been shown that, by careful protection of the eye, its disorganisation can be, if not entirely prevented, at least considerably postponed. In sum, when we consider how difficult it must be to permanently and perfectly protect the eyeball of an animal from external irritation, and, moreover, that even then we have not excluded the possible participation of a vascular factor, we are driven to admit that the direct trophic action of nerve, tried by the test case of the fifth nerve, remains unproven. And in questions of this nature, until an action has been proved to exist, it does not exist, however probable it may appear.

**Stimuli.**—Any cause that provokes a nerve to action is called a *stimulus*. The evidence that a nerve has been stimulated is in the immense majority of cases a *movement* of some kind; each individual 'ego' may, however, experience for himself another evidence of nerve-stimulation in the form of a *sensation*, without manifesting the fact by any external movement. The evidence of sensation in other persons or in animals is furnished by the movements whereby their feelings are, or appear to be, expressed. Such evidence is indirect; the observer hears statements, or sees and interprets the significance of muscular expressions; these are, of course, movements or attitudes, and obviously the value of such evidence varies greatly with the aptitude and training of the observer. A third kind of evidence is available in the laboratory on exposed nerves; the *electrical state* of nerve is altered when it is rendered active; and the alteration (negative



variation or current of action) is demonstrable by the galvanometer or by the electrometer (pp. 308, 314, 387). Movement as evidence of the passage of motor impulses, the negative variation as evidence of the passage of centripetal or centrifugal impulses, are '*objective*' signs. Sensation, as evidence of the passage of sensory impulses, is a '*subjective*' sign. Movement, as evidence of the passage of sensory impulses, is a mixed sign—*i.e.* a movement is seen, and *inferred* to be an expression of sensation. Objective signs are less liable to fallacy than subjective or mixed signs. Hence, as a rule, it is easier to ascertain loss of motion than loss of sensation.

*Conductivity.*—Nerve, in common with all protoplasm, possesses excitability, and the salient characteristic of this excitability in the case of nerve is its transmission along the fibre; hence it is termed *conductivity*. The transmission is not, as in the case of circulation, any actual transmission of matter, but only the transmission of a state of matter from particle to particle. The direct local excitability of nerve to artificial stimuli, applied at any point of its course, is sometimes distinguished from its indirect excitability to the impulse starting from that point and transmitted onwards. The distinction has its justification in experimental and in clinical facts, for it sometimes happens that the local excitability to a direct stimulus is lost at a part of the nerve that can still be traversed by impulses transmitted from above. This is apt to occur clinically during the progress of recovery from paralysis, owing to previous lesion or disease of motor nerve; it has been observed that muscles may be voluntarily set in action through motor nerves which have not yet recovered their electrical excitability. The converse may be experimentally demonstrated, *viz.* a nerve may be rendered impermeable while still remaining directly excitable.

Stimuli may be natural or artificial. Natural stimuli may arise in the brain or at the periphery, and their consequent impulses may pass from the brain to the periphery, or from the periphery to the brain. All appreciable qualities of objects in the surrounding world are natural stimuli at the sensory periphery, all consequent volitional impulses and ideas of intended movements are natural stimuli at the supreme motor centre. The natural stimuli at the sensory periphery are the physical qualities of objects that excite *smell, sight, hearing, taste, touch*, and possibly '*muscular sense, thermic sense, and pathic sense.*'

To these qualities we give the names smell, light, sound, taste, smoothness or roughness, resistance and extension, weight, heat, cold, pain. All these stimuli affect us best when we meet them half-way by some action of our own, by a state of voluntary attention, or by actual movements. An object is best seen when it is looked for, a sound best heard when it is listened for. We generally sniff to smell, and move in our mouths substances that are to be tasted, and movement on our part is necessary for anything like delicate appreciation of the nature of a surface, or the weight and size of an unseen object.

Artificial stimuli are such as are applied experimentally. They may be *mechanical*, *chemical*, *thermic*, or *electrical*, and the latter may be in the form of *induced*, *constant*, or *static* electricity. Of these various kinds of stimuli, the electrical, in the form of induced currents, is that of most frequent and convenient experimental application. The effects of the constant current have also been exhaustively studied. Those of static electricity are only incidentally observed, this form being rarely employed in the laboratory; '*unipolar stimulation*' (*i.e.* only one pole being applied to the nerve) is due to a discharge of static electricity, and is demonstrated as a fallacy to be guarded against in experiments.

*Mechanical stimulation.*—A sudden smart blow upon a motor nerve causes muscular contraction; a blow upon a sensory nerve causes pain. This is easily experienced on the ulnar nerve, which is a mixed nerve composed of sensory and of motor fibres. When it is struck just where it lies upon the bone, there result the sensation of tingling or pain, and a twitch of the muscles that are served by the nerve. In the laboratory, mechanical stimulation is applied by means of Heidenhain's tetanomotor, or of an instrument devised for the same purpose by Tigerstedt. This form of stimulation supplies the best available means for the demonstration of electrotonic alterations of excitability in the intrapolar region of a nerve during the passage of the constant current.

Of *thermic* stimuli there is little to be said; if heat be applied *suddenly* it may act as a stimulus; also, upon a change from a medium of low to one of high temperature, nerves are prone to fall into a *tetanic* state, the muscles that they supply becoming contracted, and so remaining for long periods.

The action of *chemical* stimuli has been carefully studied, especially as regards their comparative action upon muscle and



upon nerve, and, apart from experiments in which the fallacy due to self-excitation through closure of a nerve or muscle-current could occur, we may name ammonia as acting specially upon muscle, while glycerin or common salt would be selected for the excitation of nerve; removal of water is in this case the excitant—a nerve is stimulated as it dries.

**Electrical stimulation. Effects of the constant current.**—These are studied upon nerve-muscle preparations. The nerve is laid across two unpolarisable electrodes, through which it receives the current from a battery. The effects to be studied are—(1) those that take place when the current commences and ceases to flow through the nerve, *i.e.* at *make* and at *break*; (2) *electrotonic currents*, which spread along the nerve on either side of the two electrodes; (3) *alterations of excitability*, which accompany these electrotonic currents.

**Make and break effects. Pflüger's law** —When the current is *made* or when it is *broken*, or at both these events, there is contraction of the muscle, which is evidence that the nerve has been stimulated. In other words, the constant current stimulates the nerve when it *commences* and when it *ceases* to pass, but does not stimulate the nerve *while it is passing*. If now attention be given to the *strength* of current and to its *direction*, it is found that the contractions at make and at break of the current appear or fail to appear in a regular order. This is called Pflüger's law of contractions.

The terms *ascending* and *descending* are those in common use, and signify direction of current in the nerve in relation to nerve-centre. Current is 'ascending' when its direction in the nerve is from muscle towards centre, 'descending' when its direction in the nerve is from centre towards muscle. But it simplifies matters to attend particularly to the points where the current enters and leaves the nerve; the point where the current enters is the *anode* (+), the point where the current leaves is the *kathode* (—).

The explanation of the above 'law' or formula is as follows:—As stated below, when a current commences to flow, *i.e.* at *make* of a current, excitability of the nerve is diminished at and near the anode, increased at and near the kathode. A sudden increase of excitability is equivalent to a stimulus; therefore *at make of a current the stimulus is at the kathode*. As stated in the next paragraph, when a current ceases to flow, *i.e.* at *break* of the current, excitability at and near the anode suddenly recovers up

to and beyond its normal level, while at and near the kathode it suddenly falls down to and beyond its normal. A sudden change of excitability from below normal to such normal, or above it, is equivalent to a sudden increase of excitability, and furnishes a stimulus; therefore *at break of a current the stimulus is at the anode*.

With a current of 'medium' strength there is contraction at make and break of ascending and of descending currents, contractions at make being excited from the kathode (—), contractions at break being excited from the anode (+), and the formula reads—

ASC.		DESC.	
m.	b.	m.	b.
C.	C.	C.	C.

With a 'weak' current only the more efficient stimulus is effective. The sudden increase at the kathode when the current is made, is more effectual than the sudden release at the anode when the current is broken.

Hence, with a weak current contractions appear at make only with either direction of current, and the formula reads—

ASC.		DESC.	
m.	b.	m.	b.
C.	O.	C.	O.

With a 'strong' ascending current the point of stimulation at make is the kathode, but between it and the muscle lies the anode at which excitability is diminished. With such a strong current the diminution is sufficient to block the passage of the stimulus from the kathode. Hence the make contraction fails to appear. At break, the stimulus at the anode, having no obstacle between it and the muscle, produces a contraction. With a strong descending current the anode does not separate the kathode from the muscle, and the make stimulus at the kathode produces a contraction. At break, however, the stimulus is produced at the anode, between which and the muscle there intervenes the kathode where excitability sud-

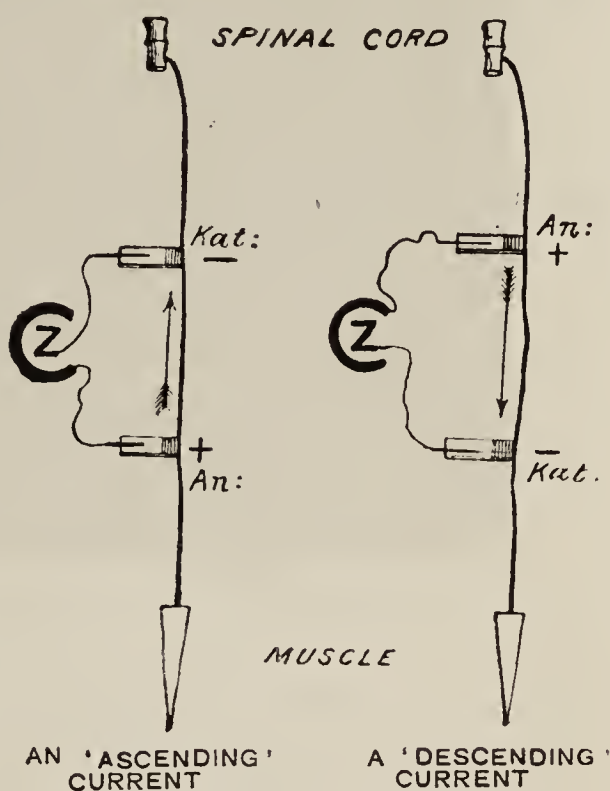


FIG. 159.



denly diminishes ; the diminution is sufficient to block the stimulus from the anode, and the formula reads—

$$\begin{array}{cc} \text{ASC.} & \text{DESC.} \\ m. & b. \\ O. & C. \end{array}$$

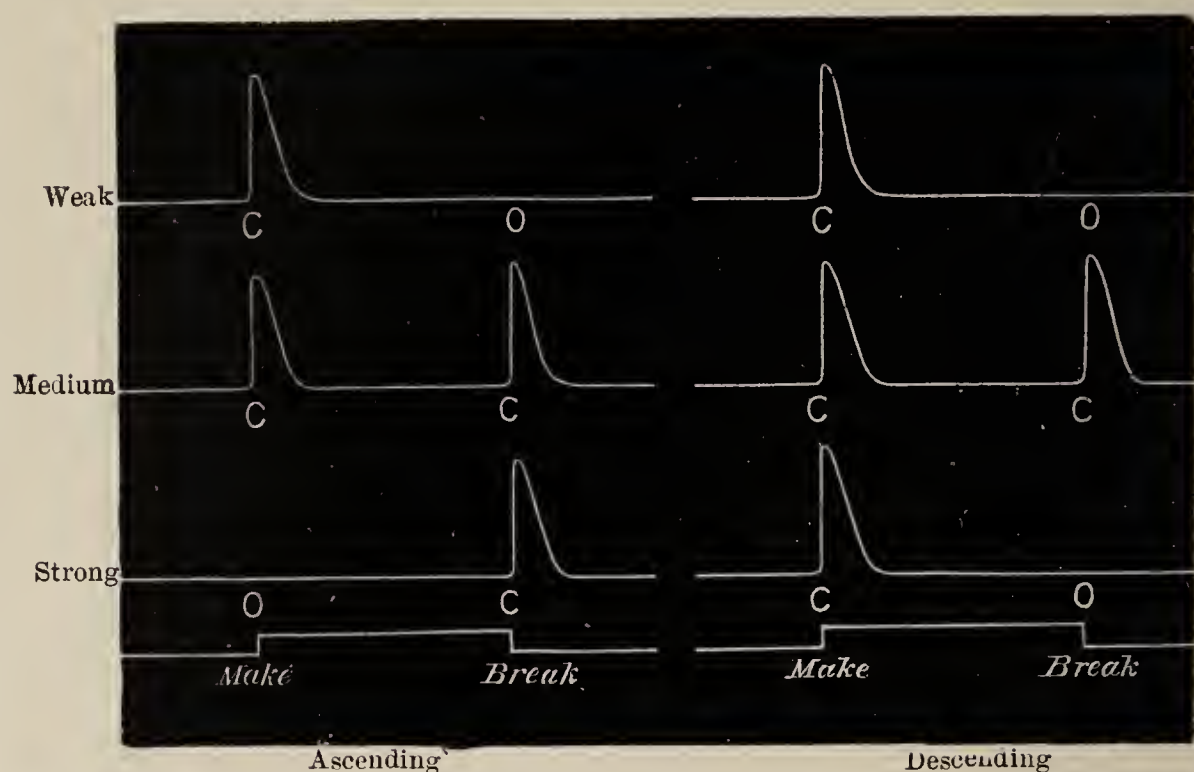


FIG. 160.—PFLÜGER'S LAW OF CONTRACTIONS ON NERVE-MUSCLE PREPARATIONS.

*Ritter's tetanus.*—It frequently happens that at break of the galvanic current the muscle enters into tetanus ; this is due to an after-anodic excitation, and the effect is also demonstrable by the galvanometer ('positive polarisation current,' p. 373).

According to recent observations by Gotch, variations of temperature alter the relative efficacy of long and of short stimuli applied to nerve, long stimuli being favoured by low, short stimuli by high temperature.

**The formula of contraction on man.**—The above statements are based upon experiments with the nerves of frogs ; experiments on man give results of which the principle is the same, but with differences that are owing to differences in the conditions of experiment. In the case of a frog's nerve cut out of the body, the points of entrance and of exit of the exciting current are usually chosen some distance apart, so that the kathode and anode are distinct, the current entering the nerve at one electrode and leaving it at the other. In the case of human nerve the conditions are different, for the nerve is imbedded in the tissues below the skin. A pair of electrodes cannot be applied to a nerve so as to send a current in at one point, out at another point ; so that it is incorrect to speak

of 'ascending' and 'descending' currents under such circumstances, and it is useless to attempt to study the effects when both electrodes are applied along the course of one and the same nerve. We must apply one electrode only to the nerve, and attend to its effects alone, completing the circuit through a second electrode, which is applied, according to convenience, to some other part of the body. Confining our attention to the first electrode, let us see what will happen according as it is *anode* or *kathode* of a galvanic current. If this electrode be the anode, current enters the nerve by a series of points, and leaves it by a second series of points; the former or proximal series of points collectively constitutes the *polar* zone or region, the latter or distal series of points collectively constitutes the *peripolar* zone or region. In such case the polar region is the seat of entrance of current into the nerve, *i.e.* is *anodic*; the peripolar region is the seat of exit of current from the nerve, *i.e.* is *kathodic*. If, on the contrary, the electrode under observation be the kathode, current enters the nerve by a series of points collectively constituting a 'peripolar' region, and it leaves the nerve by a series of points collectively constituting a 'polar' region. The current at its entrance into the body diffuses widely, and at its exit it concentrates; its 'density' is greatest close to the electrode, and the greater the distance of any point from the electrode, the less the current density at that point; hence it is obvious that the current density is greater in the polar than in the peripolar region.

These conditions having been recognised, we may apply to them the principles learned by study of frog's nerve under simpler conditions. Seeing that with either pole of the battery, whether anode or kathode, the nerve has in each case points of entrance (constituting a collective anode) and points of exit to the current (constituting a collective kathode), and admitting as proved that make excitation is kathodic, break excitation anodic, we may with a sufficiently strong current expect to obtain a contraction at make and at break with either anode or kathode applied to the nerve. And we do so in fact. When the kathode is applied, and the current is made and broken, we obtain a *kathodic make contraction* and a *kathodic break contraction*; when the anode is applied, and the current is made and broken, we obtain an *anodic make contraction* and an *anodic break contraction*. These four contractions are, however, of very different strengths; the kathodic make contraction is by far the strongest;



the kathodic break contraction is by far the weakest ; the kathodic make contraction is stronger than the anodic make contraction ; the anodic break contraction is stronger than the kathodic break contraction. Or, otherwise regarded, if, instead of comparing the contractions obtained with a ‘ sufficiently strong ’ current, we

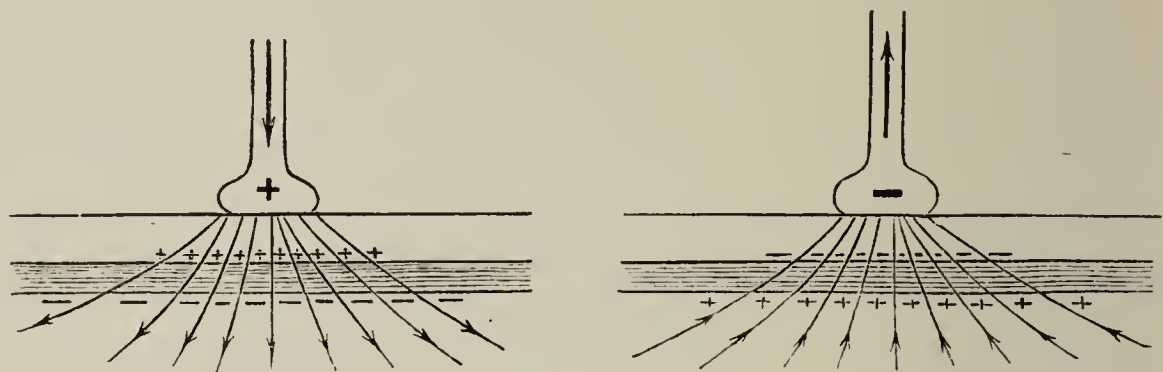


FIG. 161.

ANODE OF BATTERY.

KATHODE OF BATTERY.

Polar region of nerve is anodic.  
Peripolar region of nerve is kathodic.

Polar region of nerve is kathodic.  
Peripolar region of nerve is anodic.

observe the order of their appearance with currents gradually increased from weak to strong, we shall find that the kathodic make contraction appears first, that the kathodic break contraction appears last, and the formula of contraction for man reads as follows :—

Weak current .	.	.	K.C.C.		—		—		—
Medium current	.	.	K.C.C.		A.C.C.		A.O.C.		—
Strong current	.	.	K.C.C.		A.C.C.		A.O.C.		K.O.C. <sup>1</sup>

That such should be the normal order of appearance is fully accounted for by the following considerations :—

In the	The nature of the stimulus is	The situation of stimulus is	Conditions of excitation
K.C.C.	Kathodic	Polar	= best stimulus in best region
A.C.C.	Kathodic	Peripolar	= best stimulus in worst region
A.O.C.	Anodic	Polar	= worst stimulus in best region
K.O.C.	Anodic	Peripolar	= worst stimulus in worst region

which also account for an apparent anomaly, viz. that sometimes the anodic closure contraction precedes the anodic opening contraction, while sometimes this order is reversed ; this difference depends upon relative current densities in the two regions, which are determined by the nature of the tissues by which the nerve is surrounded. In testing this point we shall hardly fail to

<sup>1</sup> K.C.C. = Kathodic closure contraction.    A.O.C. = Anodic opening contraction.  
A.C.C. = Anodic closure contraction.    K.O.C. = Kathodic opening contraction.

notice a very evident token of the fact that polar and peripolar excitations are differently localised in the nerve; the muscles that contract to anodic make are not the same as those that contract to anodic break, if, *e.g.*, the exploring electrode is applied to the median nerve. This point need not, however, be further discussed here.

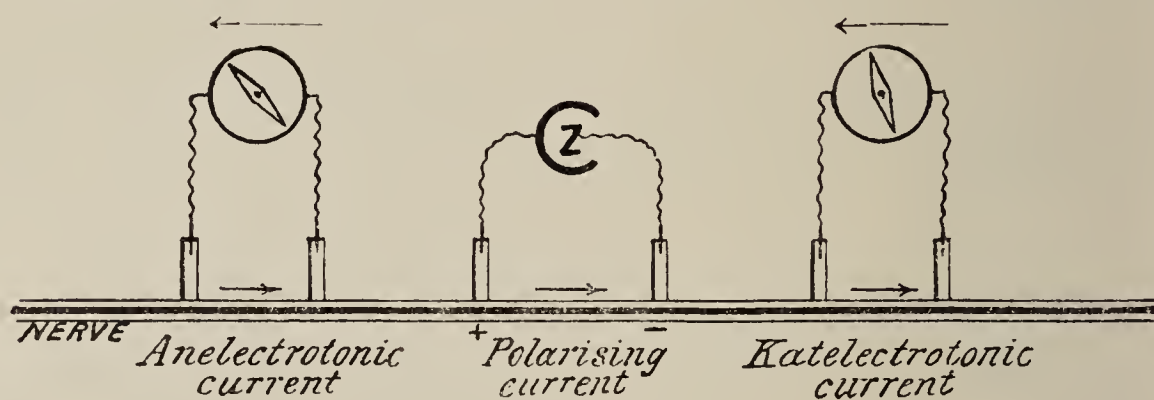
The latent period of the break contraction on man is exceedingly and constantly long ( $\cdot 05''$ ); on the frog its duration is very variable, sometimes very short, sometimes very long. With strong currents, it is usual on man to obtain tonic contraction during the passage of the current—*galvanotonus*—as well as single twitches at make and at break.

*The reaction of degeneration* is a term used to denote the reaction of diseased nerve and muscle on man. As regards *nerve*, the reaction of degeneration consists in an abolition of excitability to the constant current and to the induced current. As regards *muscle*, the reaction of degeneration consists in the abolition of excitability to the induced current, while the excitability to the constant current is exaggerated; the muscular contraction is greatly prolonged, and galvanotonus is easily produced. The normal contraction-formula given above is departed from, the most characteristic feature of this departure being a reversal of the normal order of appearance of K.C.C. and A.C.C.; normally K.C.C. appears with a weaker current than A.C.C., in a well-marked reaction of degeneration A.C.C. appears with a weaker current than K.C.C. There is no satisfactory explanation to be given of this reversal.

**Electrotonus.**—The term electrotonus is used to denote two distinct series of effects that are produced when the constant current is applied to a nerve. The effects are—(1) electrical currents in the nerve beyond the part of it that is traversed by the experimental current; these currents are known as the '*electrotonic currents*,' and the phenomenon is called '*electrotonus*'; (2) alterations of the excitability of the nerve in and near the part that is traversed by the current; this phenomenon is intimately associated with the electrotonus just mentioned, and is itself commonly called '*electrotonus*'; but, seeing that these are alterations that accompany electrotonus, it is better to distinguish them by the longer expression, *electrotonic alteration of excitability*. Electrotonic currents and electrotonic alterations of excitability are to be studied with reference to the poles at and near which they are produced, and may be in either of two



opposite directions. An electrotonic current in the vicinity of the anode is called '*anelectrotonic*,' an electrotonic current in the vicinity of the kathode is called '*katelectrotonic*.' The corresponding alterations of excitability that are found are likewise called '*anelectrotonic*' and '*katelectrotonic*' according as they occur in the vicinity of the anode or of the kathode. The terms '*anelectrotonus*' and '*katelectrotonus*' are frequently used indiscriminately to denote the currents or the alterations of



excitability. It is obviously advisable to use the terms electrotonic, katelectrotonic, and anelectrotonic currents, or electrotonic, katelectrotonic and anelectrotonic alterations of excitability, in preference to the terms electrotonus, katelectrotonus, and anelectrotonus, seeing that these last have double meanings.

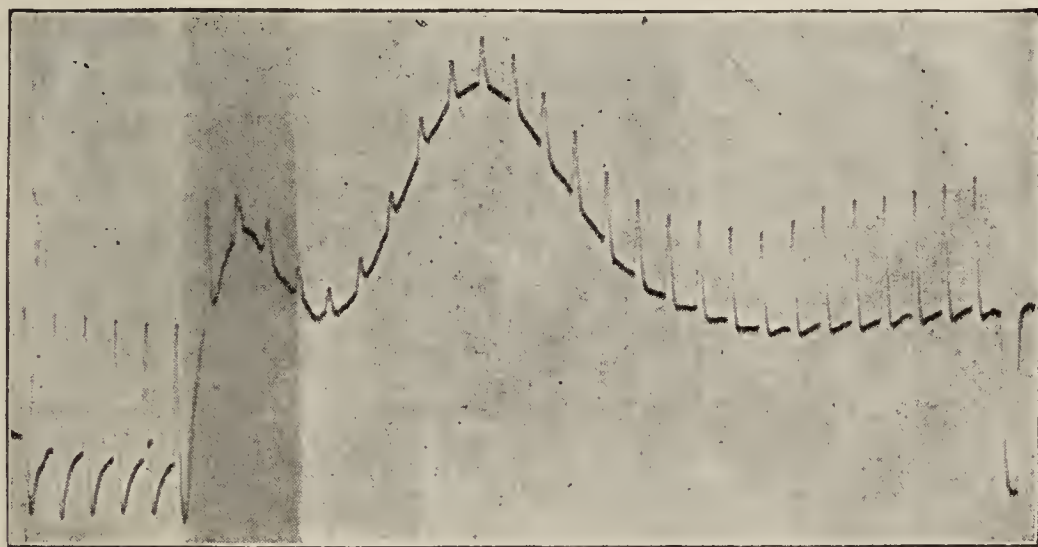
*Electrotonic currents.*—If through one pair of unpolarisable electrodes a constant current be passed into a portion of nerve, while a second pair of electrodes 'lead off' from some other portion of the nerve to a galvanometer, the galvanometer will indicate a current derived from the second pair of electrodes. This is the electrotonic current—called '*anelectrotonic*' if on the side of the anode, '*katelectrotonic*' if on that of the kathode, the former being the stronger, and their direction as indicated by the arrows in fig. 162.

An electrotonic current increases with increased strength of the polarising current, with decreased distance of the 'led-off' from the polarised region, and with increased length of the latter, and is reversed in direction by a reversed direction of the polarising current—in contrast with the '*current of action*' (p. 389), which is unaltered in magnitude by altered distance between led-off and exciting electrodes, and unaltered in direction by a reversed direction of the exciting currents. And whereas an electrotonic current bears no relation to '*current of injury*' (p. 390), but is manifested when any two points of the

longitudinal surface are led off, a current of action is generally manifested as a diminution or 'negative variation' of a current of injury between transverse section and an uninjured part of the longitudinal surface.

Electrotonic currents can be reproduced on a model composed of a core of platinum wire and a sheath of  $\text{ZnSO}_4$  solution, between the contiguous surfaces of which electrolytic polarisation gives rise to a virtual resistance, and consequent longitudinal diffusion of current in extrapolar regions. They may be provoked by induced as well as by constant currents, and this possibility must be specially borne in mind when induced currents are used for the study of the 'negative variation.'

That an electrotonic current as manifested by medullated nerve is not a purely physical phenomenon, but dependent upon the anatomical and physiological (=physico-chemical) integrity of its individual fibres (grey axis surrounded by white sheath), is, however, indicated by the following facts: 1, it is abolished by *ligature* between the polarised and the electro-

 $\text{Et}_2\text{O}$ 

0.001

FIG. 163.

Effect of ether vapour on the anelectrotonic current. A polarising current of 1 volt is made at regular intervals of 1 min. throughout the observation, and the series of anelectrotonic deflections recorded by a dead-beat galvanometer; during the time indicated by the dark bar the nerve is etherised, and the anelectrotonic deflection is temporarily much diminished. (The downward deflection at the end of the observation gives the value of 0.001 volt.) (w.)

tonic regions; 2, it is absent, or at least diminished, on *dead* and on *degenerated* nerve; 3, it is absent from tendon or a wet thread; 4, it is practically inappreciable in muscle and in non-medullated nerve; 5, it is temporarily suppressed by anæsthetics.

Admitting that electrotonic currents are due to core-polarisa-



tion, and that medullated nerve is the only tissue distinctly exhibiting the phenomenon, its probable seat of origin must be placed at the boundary between grey axis and white sheath as illustrated in fig. 164.

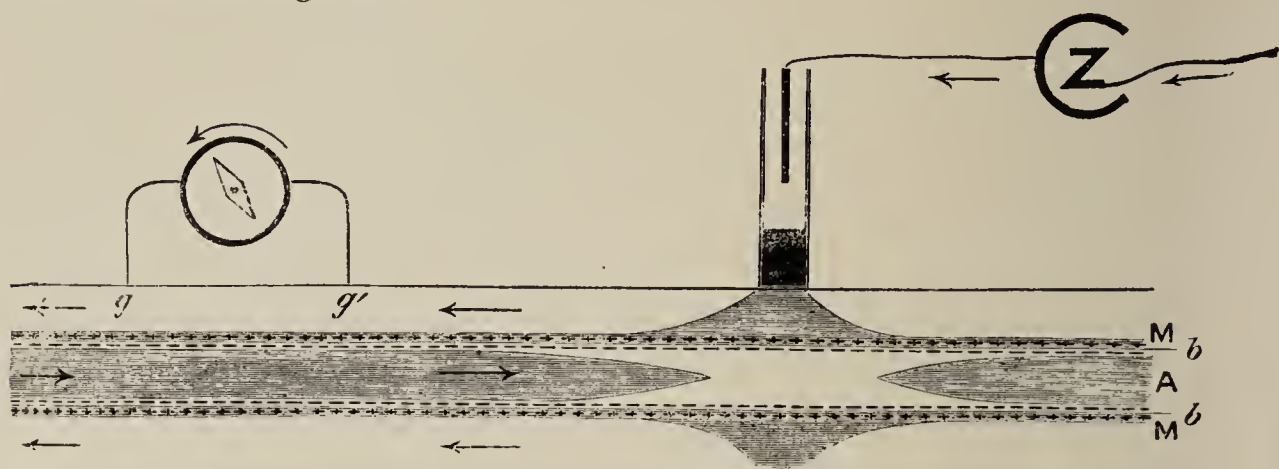


FIG. 164.—TO ILLUSTRATE ELECTROTONIC DIFFUSION IN A MEDULLATED NERVE-FIBRE.

A = axis-cylinder. M, M = medullary sheath.  $b, b$  = boundary between axis and sheath at which polarisation is supposed to give rise to virtual resistance and longitudinal current-diffusion. As represented in the figure (polarising anode and anelectrotonic current led off at  $g, g'$ ), the white side of the boundary is anodic, the grey side of the boundary is kathodic, in each case in diminishing density with increasing distance from the polarising electrode.

The after-electrotonic currents obtained when the polarising current is cut off are as follows:—In the intrapolar region the after-current is in the opposite direction, unless the polarising current has been 'strong' and of short duration, when an after-current in the same direction is obtained (du Bois-Reymond's 'positive' polarisation current). In the extrapolar region the after-current on the side of the kathode is in the same direction as the katelectrotonic current; the current on the side of the anode is at first in the same, subsequently in an opposite, direction to that of the anelectrotonic current.

Du Bois-Reymond's positive polarisation current is demonstrable on nerve and on muscle. Strong currents of short dura-

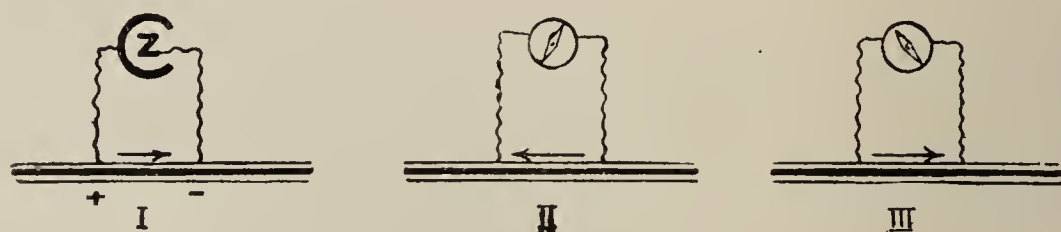


FIG. 165.—INTRAPOLAR AFTER-CURRENTS.

- I. Original, or polarising current.
- II. Ordinary polarisation after-current (negative).
- III. 'Positive' polarisation after-current.

tion are favourable to its manifestation, and the nerve or muscle must be alive. With weaker currents of longer duration, ordi-

nary polarisation currents, negative to the original current, are obtained, and these effects are obtainable on dead as well as on living muscle or nerve. Hermann has shown that the 'positive' polarisation current is in reality an action current arising from an after-anodic excitation.

*Negative variation of electrotonic currents. Positive variation of polarising currents, or polarisation increment.*—If a nerve through a portion of which a polarising current is passing, giving

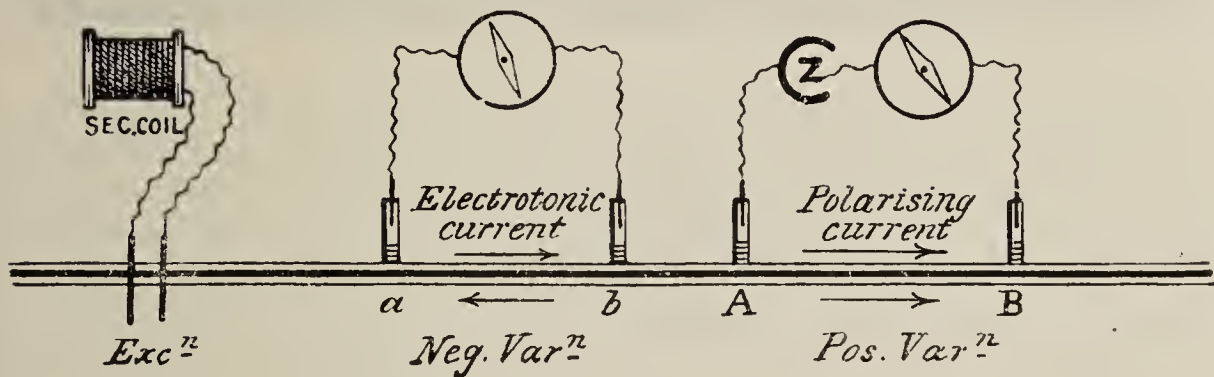


FIG. 166.

The polarising current passes in the nerve from A to B. An electrotonic current passes in the nerve in the direction *a* to *b*. During excitation the polarising current is increased, the electrotonic current is diminished. The variations are in reality 'action currents' arising as follows: During polarisation the excitability is depressed at and near the anode, viz. at A and *b*; the excitatory change sweeping down the nerve causes a larger electrical change (= a greater negativity of action) at A than at B, and at *b* than at *a*, viz. current in the nerve from A to B and from *b* to *a*.

rise to electrotonic currents in the remainder of the nerve, be excited by faradisation, the strength of the polarising and of the electrotonic currents will be altered; the strength of the polarising current will be increased, that of the electrotonic current will be diminished.

*The paradoxical contraction.*—If the branch B C of a *divided* nerve be stimulated at C by an induced or constant current, the muscle M contracts. This is not a reflex contraction, seeing that the nerve is separated from the spinal cord. It is due to an electrotonic current spreading along the fibres of B C beyond the point of junction B, and exciting in the joint portion A B adjacent fibres that are distributed to the muscle M.



FIG. 167.—PARADOXICAL CONTRACTION.

*Electrotonic alterations of excitability.*—If, while a polarising current is passed through a portion of nerve in the manner above described, the excitability be tested at various parts of the nerve, differences will be found—namely, an *increase* of excitability near the *kathode*, a *diminu-*



tion of excitability near the *anode*. Or otherwise, if, while tetanising currents are being passed through a nerve, a polarising

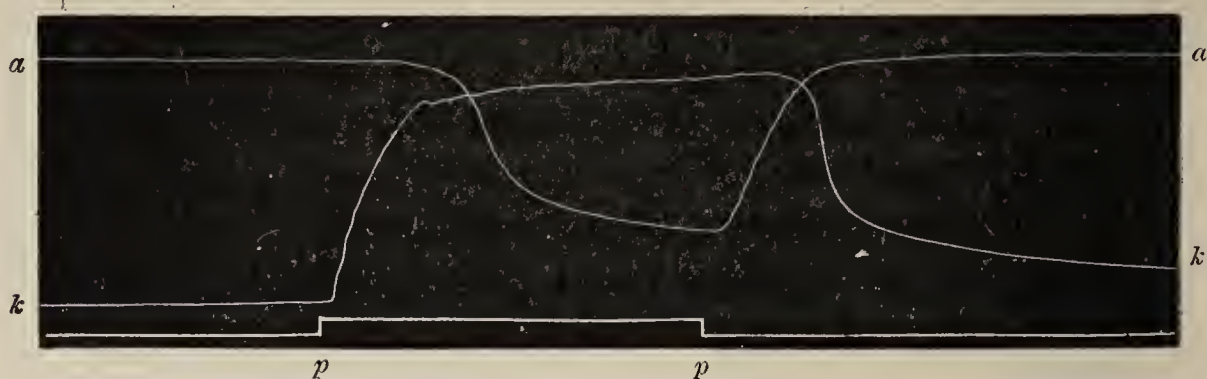


FIG. 168.—ELECTROTONIC ALTERATIONS OF EXCITABILITY. NERVE-MUSCLE PREPARATION.

Insufficient tetanising stimuli produce a tetanus (line *k k*) during the passage of a polarising current with the kathode near the exciting electrodes. A tetanus (line *a a*) is cut down during the passage of a polarising current during the period *p p* with the anode near the exciting electrodes.

current is passed through the nerve for a given period, it is found that the effect of the tetanising current is modified during that period—*increased* if the polarising *kathode* be near the point of application of the tetanising current, *diminished* if the pola-

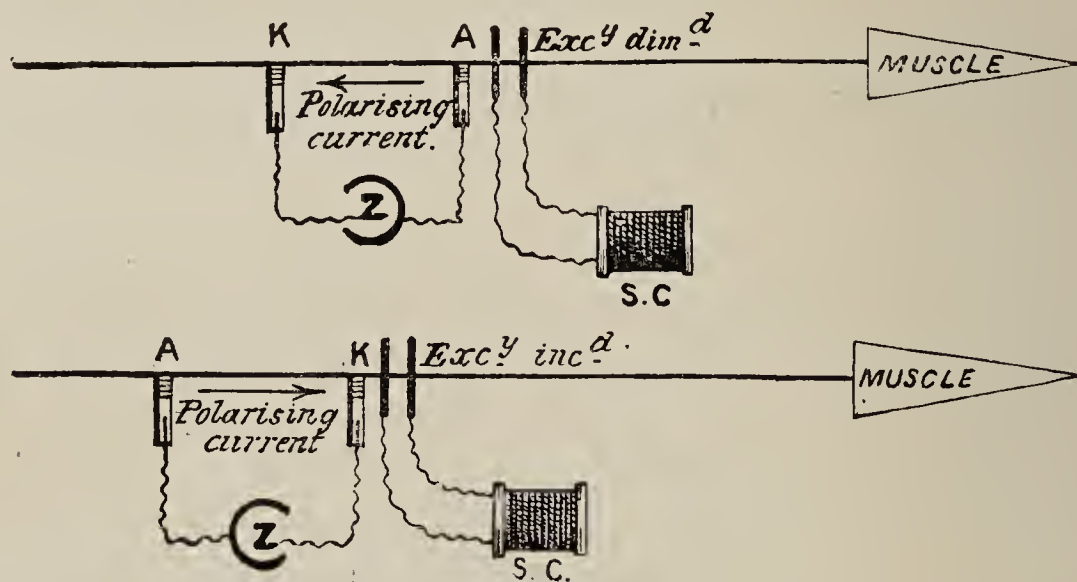


FIG. 169.

rising *anode* be near the point of application of the tetanising current, and is less and less farther and farther from the anode. The kathodic augmentation and the anodic diminution are greater and smaller according as the distance between the polarising and exciting electrodes is smaller and greater; this is expressed in the following diagram (fig. 170).

**Experiments on man (w.).**—The above conclusions are based upon experiments made upon frog's nerves. Experiments on man give similar results, with minor differences owing to differences in the conditions of experiments. As regards frogs' nerve, it is isolated, and the test is applied separately from the pola-

rising current, induction currents being most convenient for testing the extrapolar region, while mechanical stimuli are best adapted for testing the intrapolar region. As regards human nerve imbedded in the tissues, such a mode of testing is not pos-

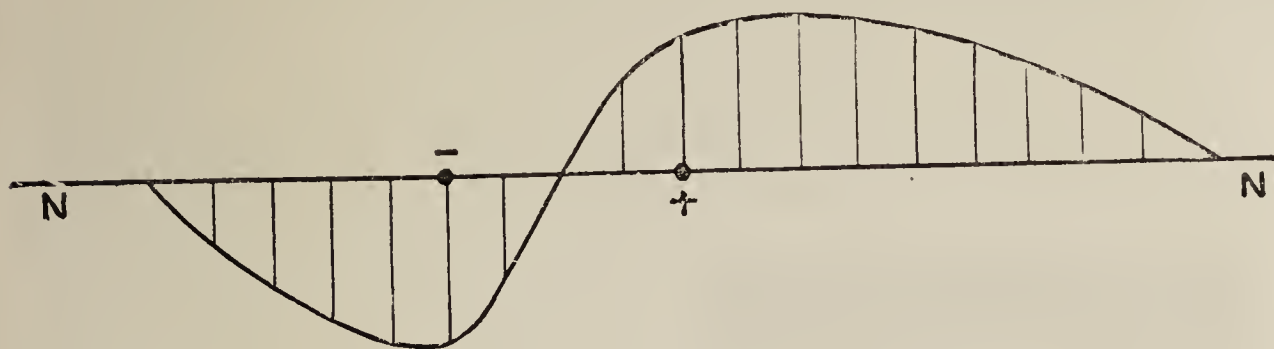


FIG. 170.

NN is a line to denote a nerve and at the same time the level of normal excitability, *i.e.* that observed in the absence of any polarising current; + and - denote respectively the anode and kathode of a polarising current; the region included between them is the intrapolar region. The degree of altered excitability at various points of the nerve near the anode and kathode respectively is indicated by perpendicular lines below and above the normal NN. A line joining their summits indicates the mode of distribution of the altered excitability at and near the anode and kathode of the polarising current. The excitability of the anode is most diminished at the anode, less diminished in its immediate vicinity, still less diminished at more remote points. The excitability of the kathode is most increased at the kathode, less increased in its immediate vicinity, still less increased at more remote points (Pflüger).

sible, and it is necessary to adopt some means for ensuring that the test shall coincide with the polarised region of the nerve. This can be effected by conjoining in one circuit the testing with the polarising current, or in the case of mechanical stimuli by applying the latter through the electrode of the polarising current.

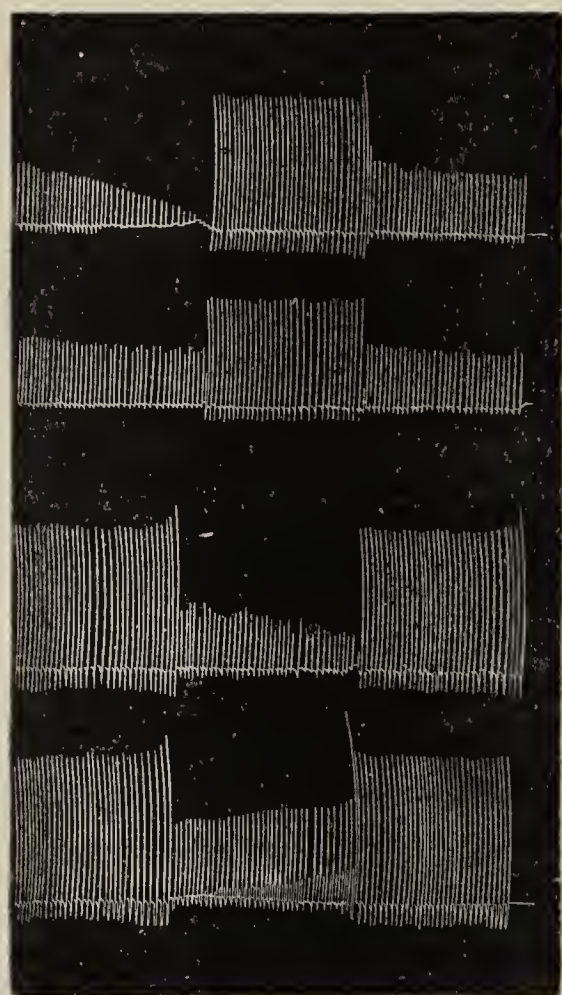
The effect of an induction shock alone is compared with its effect in the presence of a polarising current; the difference observed is owing to the modification of excitability produced by the polarising current. The polar effects under the conditions of application of electricity to the human body, and the distinction between polar and peripolar excitation (p. 368), must be borne in mind in this connection. Remembering that a current entering or leaving the body through an electrode applied over a nerve has in that nerve a polar and a peripolar region—polar of the same sign as the electrode, peripolar of the opposite sign; remembering, further, that make excitation is kathodic, break excitation anodic, it is easy to understand the effects of various possible combinations between testing and polarising currents. If the induction current is used as the test, the combinations



that it is possible to form at the exploratory electrode with a polarising current in the same circuit are as follows :—

1. Kathode of testing current, and of polarising current.
2. Anode of testing current, and of polarising current.
3. Kathode of testing current, and anode of polarising current.
4. Anode of testing current, and kathode of polarising current.

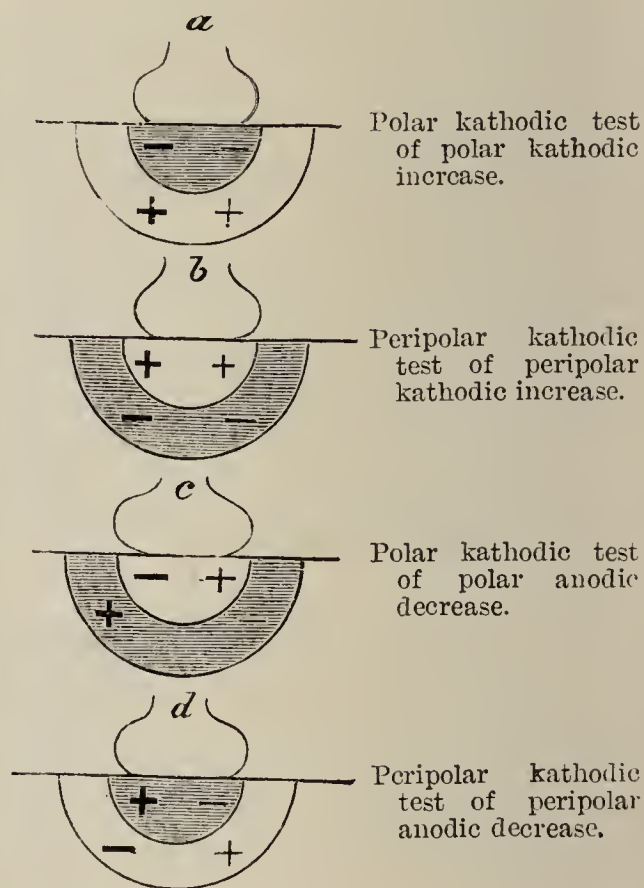
Putting these four cases to the test, it will be found that excitability is *increased* in the polar region when it is *kathodic*,



After      ←      During      Before  
                 polarisation

FIG. 171.

Electrotonic alterations of excitability of human nerve.



*diminished* when it is *anodic*, and that similarly, but in smaller degree, excitability is increased in a peripolar kathodic region, diminished in a peripolar anodic region.

Similar results follow the application of other tests—make and break of a constant current, alone, and in the presence of a polarising current in the same circuit—mechanical stimulation, alone, and during the passage of a polarising current—viz. increased excitability in a kathodic region, diminished excitability in an anodic region.

*Sensory nerves.*—The statement applies to motor and sensory

nerves alike. On the former it may be demonstrated by the graphic method, on the latter it requires to be experienced in the form of sensation by the observer himself becoming the subject of experiment. Experiments on man present the advantage that alterations of sensory excitation can be better appreciated by the subjective evidence of sensibility than by the objective signs from which inferences are drawn in the case of animals.

*After-effects.*—Not only does the constant current cause a modification of excitability during its passage, but it leaves excitability altered after it has ceased to pass. The ultimate after-effect is always an increase of excitability, but in the case of the cathodic after-effect the increase is preceded by a brief period of diminished excitability.

**Rapidity of transmission of nervous impulses.**—The rate at which an impulse travels along a nerve can best be determined for *motor transmission* along an ordinary mixed nerve. The experiment is as follows:—



FIG. 172.

An induction shock is sent into the nerve at B as close as possible to the muscle, and the interval between the application of the stimulus and the commencement of the resulting contraction is recorded. A second induction shock is sent into the nerve at A as far as possible from the muscle; the interval between stimulus and contraction is measured as before. This second interval is longer than the first by a fraction of a second, and the difference is owing to the time that the stimulus at A requires for its passage from A to B. Given this difference of time and the distance between A and B, it is easy to calculate the rapidity per second. Thus,

if the difference is  $\frac{1}{100}$  sec., and the distance  $\frac{1}{2}$  meter, the rapidity is 50 meters per second. This is about the rapidity of motor transmission in *human nerve*, as may easily be determined



FIG. 173.—VELOCITY OF MOTOR IMPULSE IN HUMAN NERVE.

by recording a pair of contractions of the flexor muscles of the fingers—the one contraction caused by an induction shock to the cervico-brachial plexus above the clavicle, and a second



contraction by an induction shock to the median nerve at the bend of the elbow.

The experiment was first made *on the frog* by Helmholtz, just after it had been proclaimed by Johannes Müller that the rapidity with which nervous impulses travel must remain concealed from us. Helmholtz found that at ordinary temperature the rapidity is about 30 meters per second, and he soon afterwards made measurements on man with the same result, viz. a rapidity of 30 meters. Small animals are obviously less adapted to such determinations than large; the rapidity is not markedly lower on cold-blooded animals at ordinary temperature, but is greatly diminished at low temperatures. On the frog, for instance, it is found to be only 1 meter per second at 0°.

The rate of transmission of the electrical disturbance (*negative variation* or current of action, see p. 389), which accompanies the nerve-impulse, has been measured by Bernstein with the same result, viz. a rapidity of about 30 meters per second. It need hardly be added that neither a nervous impulse nor its accompanying electrical change is of the same order as the transmission of electricity along a conductor; the velocities of the two processes are respectively 30 and 300,000,000 meters per sec. The nerve-impulse implies the transmission of a physiological state, and is similar to the excitatory wave in muscle; this last has a lower velocity, viz. in voluntary muscle 3 m., in the contractile tissue of medusa 0.5 m., in frog's cardiac muscle 0.1 m., in the intestine and in the ureter only 0.025 meter.

Attempts have been made to determine the rapidity of transmission of impulses *along sensory nerves*, and it is stated to be the same as that along motor nerves. The method followed has been to stimulate at points of nerve near and far from a centre, and to record the interval between each stimulus, and the consequent reflex contraction. The difference is attributed to time occupied in the passage of a centripetal impulse along the afferent channels. Obviously, however, the data towards this conclusion are less cogent than in the case for motor transmission, for the impulse has to pass a nerve-centre, and the delay which it there undergoes is not only great but *variable*. The data that are obtained by stimulating at points of the skin near and far from the cerebro-spinal axis (*e.g.* at the shoulder and at the fingers, or at the thigh and at the foot), and comparing the reaction times of voluntary response to such stimuli, are still less satisfactory

for the purpose of estimating rate of transmission in sensory nerves. The variants in such reaction times are too great, both as regards central delay and cutaneous sensibility, to permit of exact data being obtained.

The only exact method of measuring the rate of a sensory impulse would be to measure the rate of propagation of the negative variation along a pure afferent nerve. This has not been done, and would, indeed, be almost superfluous, for we may legitimately accept in its stead the rate of transmission of the negative variation in mixed nerves as an index of the speed at which an impulse travels along all medullated fibres in either direction.

It is probable that the rate of transmission is lower in non-medullated than in medullated fibres; but there are no exact data in point.

**Conduction in both directions.**—Afferent nerves normally conduct impulses in one direction only—from periphery to centre. Efferent nerves normally conduct impulses in one direction only—from centre to periphery. And when an afferent or an efferent nerve is stimulated at some point of its course, the effect is manifested in the former case at a sensory centre, in the latter case at the motor periphery. There is, however, evidence to show that nerves of either kind are capable of transmitting impulses in both directions, though in one of these directions in each case no effect is manifest. If a pure motor nerve is stimulated at the point B, the muscle contracts, and gives evidence that a



FIG. 174.

nervous impulse generated at B has passed from B to C. It does, however, also pass from B to A, but to prove this requires a special test, for the motor nerve does not possess any means at the centre of revealing the fact. The electrical disturbance or 'current of action,' which accompanies any action in nerve, furnishes this proof—when B is stimulated, it is demonstrable between B and A as well as between B and C, *i.e.* it is transmitted in both directions. Similar proof has been given of the passage of impulses in both directions along afferent nerves.

A second proof as regards afferent nerves is offered by an experiment of Paul Bert. He grafted the tip of one rat's tail to



the nose or back of another rat, and when union had been effected amputated the tail near its base. The second rat was thus provided with a trunk-like appendage, and after a time obtained sensory connection with the artificial organ, and gave

evidence of sensation if it were pinched. Under these circumstances the impulse passed from base to tip of the tail, whereas formerly it passed from tip to base.

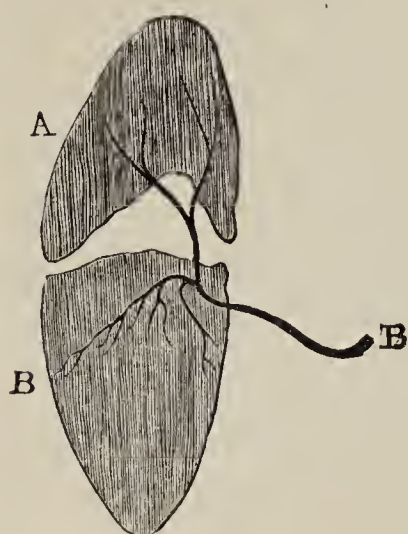


FIG. 175.

Kühne's gracilis experiment, demonstrative of double conduction in efferent nerves. The muscle is divided at its tendinous intersection into two parts, A and B; excitation localised at the peripheral end of the nerve to B, causes contraction of A as well as of B.

Kühne's gracilis experiment proves the point on an efferent nerve. The gracilis muscle of the frog is in two portions with a tendinous intersection, and supplied by a nerve that divides into two chief bundles; excitation strictly limited to *one* of these bundles causes *both* portions of the muscle to contract.

**Inequalities of excitability.**—A nerve is not equally excitable at all points of its course, nor in all the fibres of which it is composed. Instances in point are—(1) the comparative excitability of the motor nerve-fibres distributed to extensor muscles of a limb, and of those distributed to the flexor muscles, and (2)

the comparative excitability of the vaso-dilatator and vaso-constrictor fibres contained in a mixed nerve. If the sciatic nerve of a frog be stimulated with tetanising currents of gradually increasing strength, the foot will at first become flexed, and, as the current strength increases, the flexion will give place to extension. This is considered to show that the nerve-fibres to flexor muscles are excitable by weaker stimuli—*i.e.* are more excitable than the nerve fibres to extensor muscles (Ritter, Rollett). Analogous observations have been recorded, relating to the excitation of vasomotor nerves; and the usual result of primary vaso-constriction followed by secondary vaso-dilatation, also the fact that with weak currents vaso-constrictor action is better manifested, while with strong currents vaso-dilatator action is more prominent, have been explained as being due to greater excitability of vaso-constrictor in comparison with vaso-dilatator fibres. But the experimental data are not very precise nor very easily obtained.

Unequal excitability at different points along the course of a

nerve can only be observed on excised nerve (for apparent inequalities in nerves imbedded in the tissues need not at all be inequalities of excitability; they are due to inequalities of excitation, according as the nerve is more or less accessible to the application of currents), and was for long an unexplained phenomenon. But it is now understood to be owing to local injury of the trunk itself, or of the large branches that must be cut through when the nerve is excised, and it is found that when a nerve is excised with the greatest care, the maxima of excitability correspond with the points where large branches come off from the trunk of the nerve.

If a mixed nerve, left in connection with centre and with periphery, be stimulated by induction shocks of progressively increasing strength, it is found that a direct effect of nerve on muscle is obtained, with a weaker stimulus than a reflex effect; this is so, even when the excitability of the cord is exaggerated by strychnia. To infer from this that efferent are more excitable than afferent fibres is not admissible; all we may justly say is that to discharge the train—afferent fibre, central cell, efferent fibre-muscle, nerve—requires a stronger stimulus than to discharge efferent fibre-muscle. Or, otherwise regarded, the spinal cell—although acting as a reinforcing relay towards an effective stimulus, and giving a more general and massive muscular excitement—opposes resistance to stimuli below a certain level of efficacy.

A notable theory, known as *the avalanche theory*, was advanced by Pflüger; it was to the effect that a stimulus gathers strength in its passage down a motor nerve, so that a weak stimulus of the nerve far from the muscle would produce a greater effect than a similar stimulus nearer to the muscle. But the fact and the theory have been discredited by subsequent observers, and it is now accepted that the nervous impulse neither gains nor loses in strength in its passage, but remains throughout of uniform strength.

*Ritter-Valli 'law.'*—The Ritter-Valli 'law,' so-called, is a simple assertion of fact, and an additional example of unequal excitability at various points along the course of a nerve. The nerve of a nerve-muscle preparation immediately after excision is more excitable at its upper portion near the cut end than at its lower portion near the muscle. With lapse of time it is found that the increased excitability of the upper as compared with the lower end gives place to diminished excitability, which gradually



falls to zero, and this loss of excitability progresses in a centrifugal manner along the nerve towards the muscle, so that, as time passes, the points A, B, C, D, lose their excitability in the order named. This statement holds good for excised *motor* nerves, and for divided motor nerves in the body, and for undivided motor nerves at death. The further statement has been

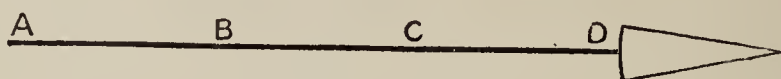


FIG. 176.

made that *sensory* or afferent nerves progressively lose their excitability in a reverse direction, *i.e.* from peripheral part first in a centripetal direction to central part last. The statement is purely conjectural, and not justified by experimental facts; it is not possible to test the point after death, and clinical experience teaches that the central end of a divided nerve can preserve its excitability for years.

**Fatigue of nerve?**—‘Fatigue’—the consequence of previous exertion of force—is caused by an actual expenditure of material and accumulation of waste product, and has as its functional characteristic a diminished capability for renewed exertion. That muscle is a force-producing organ is recognised by the

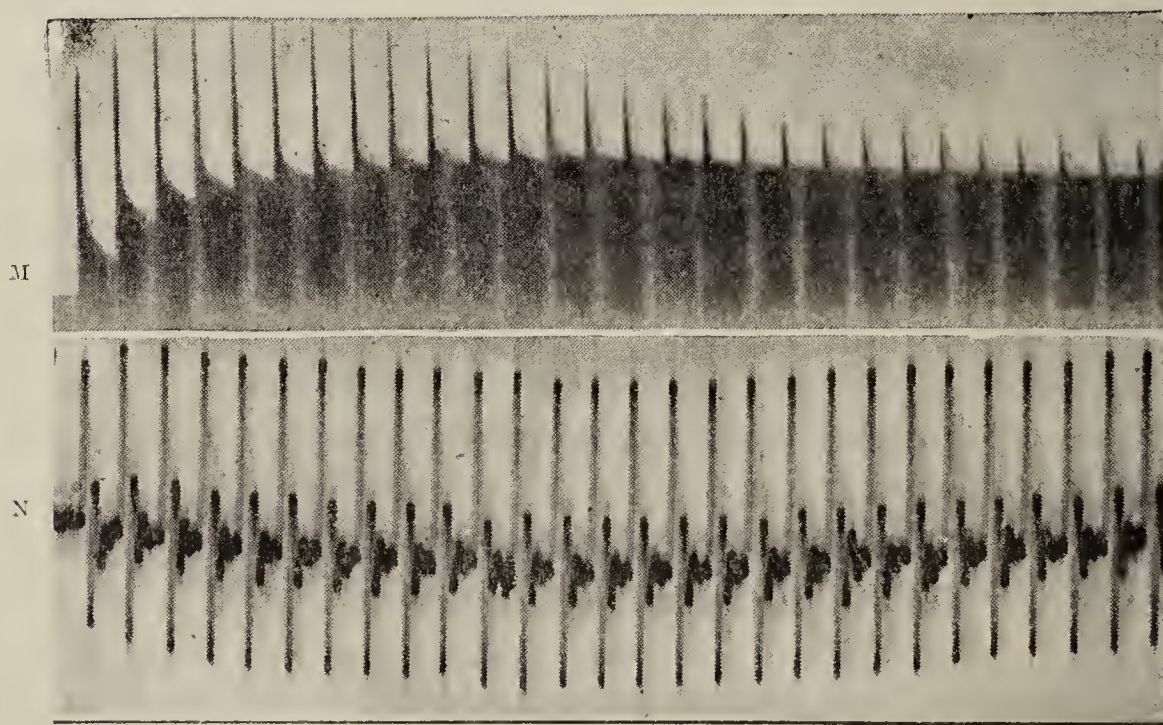


FIG. 177.

Simultaneous record of muscular contraction (upper line) and of negative variation of nerve (lower line) (W.) N.B.—The connections of the nerve with the galvanometer were such as to give the neg. var. reading upwards; the excitation of the nerve occurs at 1 min. intervals.

mechanical, thermic, chemical, and electrical changes that accompany contraction, and we have already recognised the signs of

fatigue on excised muscle subjected to direct excitation (p. 333). The case of nerve is otherwise; there is no mechanical change with its activity, nor has any direct demonstration been given of thermal or chemical changes. The only clear sign of activity in nerve itself is the 'current of action,' and this sign, while it affords evidence of some chemical change (in all probability a production of  $\text{CO}_2$ ), exhibits an almost unlimited endurance. Nerve is chiefly a force-conducting organ, and in very slight degree as compared with muscle, a force-evolving organ. Whereas muscle can without difficulty be experimentally fatigued and exhausted, nerve is practically inexhaustible; it wears extremely well—less perhaps because it does not suffer change than because any change is made good as fast as it occurs (w.)

Diminishing muscular contraction consequent upon prolonged stimulation of the nerve of a nerve-muscle preparation is no proof of fatigue of nerve. In Bernstein's experiment two nerves,  $N_1$  and  $N_2$ , are equally strongly tetanised for several minutes, but one of them,  $N_2$ , is prevented from acting on its muscle. The first nerve-muscle preparation becomes fatigued and ultimately exhausted, but that such exhaustion is not of the nerve is at once shown by unblocking  $N_2$ , which, although it has been excited as long as  $N_1$ , retains full power over its muscle. Similarly a curarised cat, after a sciatic nerve has been kept under excitation for hours, exhibits muscular contractions from the excited nerve as the curare effect wears off (Bowditch).

It must not, however, be concluded that the muscle itself is exhausted when stimulation of its nerve no longer causes it to contract. A direct stimulus to the muscle shows at once that it is still excitable. We have just seen by experiment that the nerve is still excitable. Seeing that stimulation of an excitable nerve fails to cause contraction of an excitable muscle, the conclusion follows that it is at the junction between nerve and muscle (the motor end-plate) that fatigue and exhaustion first develop (w.) The actual nature of the feeling

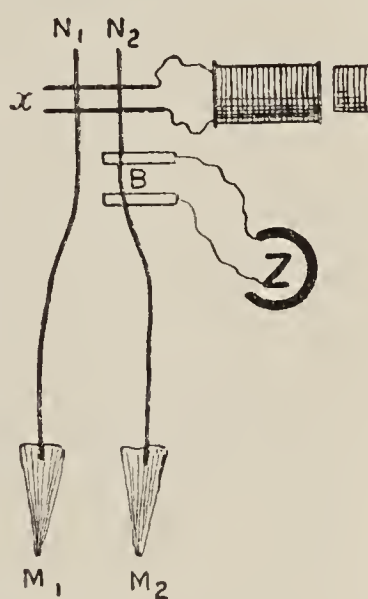


FIG. 178.—BERNSTEIN'S EXPERIMENT.

Both nerves stimulated through one pair of electrodes,  $x$ ; impulses in  $N_2$  blocked at  $B$  by a constant current. When  $M_1$  is completely exhausted,  $N_2$  is unblocked by breaking the constant current;  $M_2$  then begins to respond, *i.e.*  $N_2$  is not exhausted.



of fatigue, and the relative shares in it assumed by peripheral and by central changes, will be more suitably discussed in connection with cerebral function (p. 569).

*Inhibitory stimulation.*—We have seen that closure of a galvanic current produces a motor or excitatory effect at the kathode and an anti-motor or depressant effect at the anode. Hering regards this as an instance of the theory according to which a nerve or other excitable tissue is susceptible of two opposite changes, viz. disassimilation together with greater action or excitation, assimilation together with lesser action or inhibition. Many facts have recently been brought to light in this connection, showing that under certain conditions the stimulation of efferent nerves may be anti-motor or inhibitory as regards muscular contraction, yet hardly sufficient to justify the assumption that there are inhibitory fibres to a skeletal muscle analogous with the well-known inhibitory fibres to cardiac muscle. Richet and Biedermann have shown that the contracted muscles of the crayfish claw and the veratrinised frog's sartorius can be made to relax by tetanisation of their nerves; Kaiser has shown that a muscle put into tonic contraction by exciting its nerve with glycerin is made to relax by tetanising the nerve; and Wedensky has found that an ordinary nerve-muscle preparation may be made either to contract or to relax according to the strength and frequency of the induction shocks applied to the nerve. In all these cases we must suppose that 'interference of stimuli' (or more properly interference of excitatory states) takes place in the muscle or in the end-plate, without invoking the existence of a special class of 'musculo-inhibitory' nerve-fibres. It is obvious that only muscle that is relaxed or not fully contracted can be excited to contract, and that before a muscle can relax it must be in contraction; the same stimulus along the same nerve-fibre can therefore presumably put a resting muscle into action or an acting muscle into rest; and, indeed, according to Fick and Gad, some such double process occurs in every muscular twitch, *i.e.* in each contraction and decontraction. The *voluntary* inhibition of movement has been already alluded to (p. 351) as having its seat of consummation in the central organ.

## CHAPTER XI

\**ANIMAL ELECTRICITY*

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393	The frog's heart—The mammalian heart—The human heart.
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'ANIMAL electricity' may be regarded as having taken origin from the observation by Galvani of the spasms occurring in frogs' legs that were suspended by copper hooks to an iron balustrade (1790). We now know that such spasms are not evidence of animal electricity, but caused by currents arising from dissimilar metals. A long controversy between Galvani and Volta, supported by numberless experiments from the roughest to the most precise, a new and delicate instrument—the galvanometer—the outcome of discoveries in the collateral subject of magnetism, were, however, required before any clear understanding was reached. From the outset Galvani asserted the existence of animal currents, and denied that of metal currents; Volta denied the existence of animal currents, and asserted that of metal currents. We now know that both assertions are correct, both denials incorrect. Animal currents exist, metal currents exist; the latter in Volta's hands were the seed out of which have sprung the battery, the telegraph, the electric motor, the electric light, and the telephone; the former in Galvani's hands is the parent of the more hidden offspring, animal electricity, or the electricity evolved by living tissues.

It will be useful to repeat, as may easily be done, the chief landmark experiments to which appeal has been made during the



growth of our knowledge of animal electricity, and by which certain simple conclusions have been isolated.

1. *Galvani's first experiment (with metals)*.—Pith a frog and remove the skin, lay bare the long roots of the sciatic nerves, take away the intestines and other soft parts, leaving only a piece of vertebral column and the nerves, by which the legs can be suspended with a copper hook to an iron stand; tilt the stand

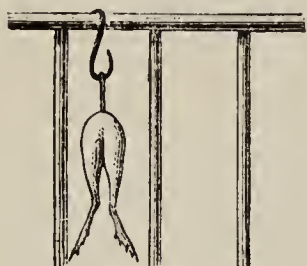


FIG. 179.—GALVANI'S  
EXPERIMENT WITH  
METALS.

until one of the legs touches the iron. Contraction occurs. This contraction is due to a completion of circuit allowing of the sudden passage of a current. Galvani thought the current was from the animal, Volta objected that it was from the contacts of copper and iron. In the controversy that followed, Galvani showed that two metals were not

necessary to the success of the experiment; Volta objected that even a single metal differs at different points, and may, therefore, give a current.

2. *Galvani's second experiment (without metals)* gives, however, proof positive that animal tissues and fluids can produce and

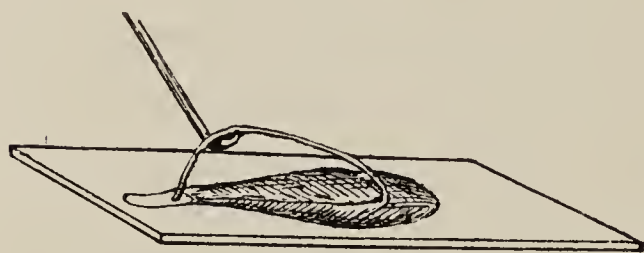


FIG. 180.—GALVANI'S EXPERIMENT WITHOUT  
METALS.

conduct a current, and is to be regarded as the foundation-stone of animal electricity. A nerve-muscle preparation is laid upon a glass plate, the nerve is raised upon a glass rod, and its freshly-cut extremity is allowed to touch

the muscle, when a contraction occurs. Here the circuit is entirely animal tissue and fluid, and the current made through it at its completion must therefore be 'animal.'

This really proved Galvani's assertion that animal currents exist, but it did not prove the non-existence of metal currents. As is now well understood, the excitation is due to 'injury-currents,' of which the circuit is suddenly completed through the nerve. Meanwhile Volta conclusively proved his assertion that metals in contact with a fluid produce a current; he invented the battery, and by his fruitful discoveries entirely overshadowed Galvani's successful demonstration.

In 1819 Oersted discovered and demonstrated the manner in which a magnetised needle is deflected by an electric current in its vicinity, and among the important consequences of the dis-

covery was the invention and construction by Nobili in 1825 of the galvanometer, which enabled subsequent observers to detect and measure the comparatively small currents derived from animal tissues.

In 1841 du Bois-Reymond began to devote himself to their study; by the invention of unpolarisable electrodes, and by his application of the method of compensation with the aid of the rheochord, he finally separated animal from metal currents, measured the former under various conditions of activity, and demonstrated the important phenomenon of the 'negative variation,' or, as it is now more usually called, the 'current of action.'

This brings us to the modern phase of the subject, in which Hermann is the chief factor; he was and is the leading opponent of du Bois-Reymond's theory that natural currents pre-exist in normal resting tissues, and is the leader of the growing school by whom animal currents are considered to be the result of chemico-physiological inequalities caused by injury or by activity.

**The electromotive properties of muscle and nerve.**—Muscle and nerve may be conveniently considered together as regards their electromotive properties, for the principles upon which the phenomena are to be explained are the same for both, and the differences—which are of a minor character—can be easily understood after the facts common to both have been mastered. These facts may be briefly formulated in three propositions:—

1. *Normal muscle (or nerve) is isoëlectric, i.e. gives no current through a galvanometer with which it is connected by a pair of unpolarisable electrodes applied to any two points of the muscle (or nerve).*

2. *Local injury disturbs this state; current is manifested through the galvanometer from the uninjured to the injured part.*

3. *Local action disturbs this state; current is manifested through the galvanometer from the resting to the active part.*

Of these three statements, the first has been the most disputed, and cannot well be completely demonstrated; the last two are easily verified.

Perfectly 'normal' muscle, *i.e.* muscle at rest and free from injury, cannot be experimented upon, because the necessary exposure is a sufficient cause of injury, and exposure is necessary because the skin, if it be not removed, itself gives current. This is true of all animals, including fishes. But that 'normal' muscle, if it could be examined, would be found to be isoëlectric or currentless, is practically certain; the heart, which can be ex-



posed without injury, is so, and muscle gives least current when it is exposed with most care, *i.e.* with least injury. Further, as the effect of local severe injury intentionally caused, a comparatively strong current is obtained, and it is highly probable that the comparatively weak currents observed from a carefully exposed muscle are due to local slight injuries incidental to exposure. The direction of current when a local injury is inflicted is best understood by comparing it with the case of the Daniell cell.

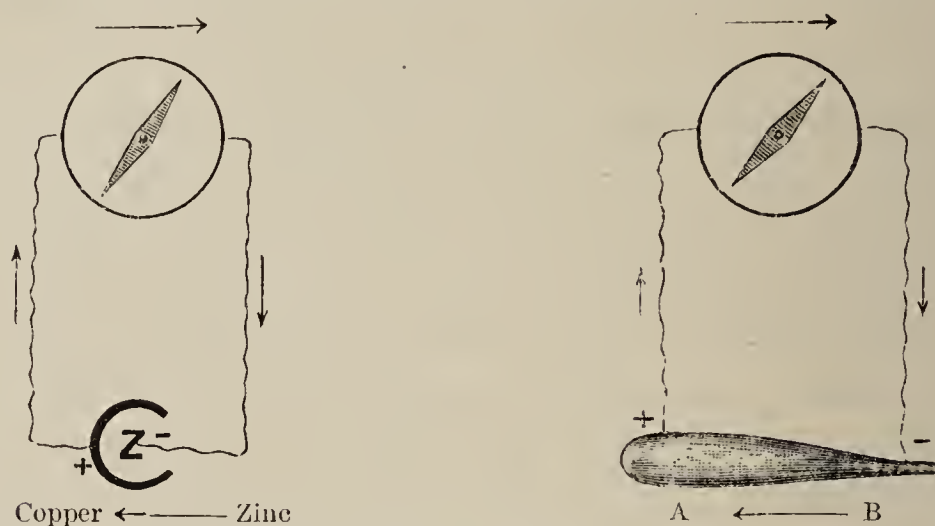


FIG. 181.

## DIRECTION OF CURRENT OF DANIELL CELL.

Through the galvanometer the current is from copper to zinc.

Through the cell the current is from zinc to copper.

## DIRECTION OF CURRENT OF INJURED MUSCLE

Through the galvanometer the current is from normal to injured part, or from resting to active part.

Through the muscle the current is from injured to normal part, or from active to resting part.

The injured muscle is thus an electromotive element comparable with the Daniell cell, and just as in the latter the zinc—at which most chemical action takes place—is the positive element, and has connected with it the negative electrode, so in the former the injured part—at which most chemical action takes place—is a positive element, and has connected with it a negative electrode. It is usual to describe direction of current with reference to its passage through the galvanometer, and to say that B, the injured part, is negative to A, the normal part. The expression ‘negativity of injury’ is sometimes used to express this relation of B to A.

A precisely similar mode of explanation is applicable to the effect of *local action*. At the seat of action most chemical change takes place; the active part is thus analogous with the zinc of the Daniell element, and the current in the galvanometer is from part at rest to part in action, in the muscle itself from part in action to part at rest. Describing direction of current with reference to its passage through the galvanometer, we say that

B, the active part, is negative to A, the resting part. The expression 'negativity of action' is sometimes used to express this relation of B to A. These considerations apply equally to muscle and to nerve.

**The negative variation of muscle (or of nerve).**—If an injured muscle giving the injury current from A to B be tetanised by

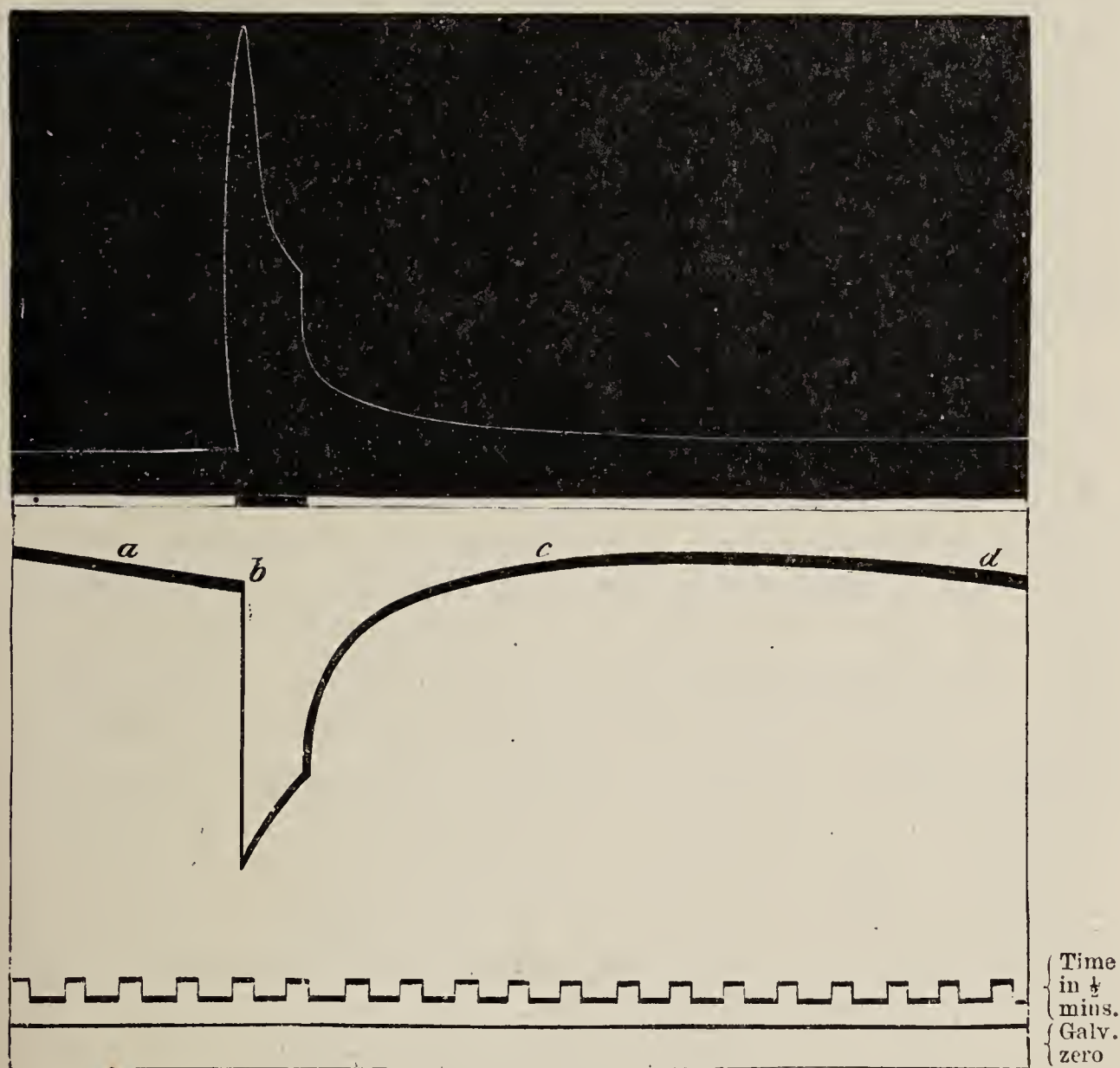


FIG. 182.—THE NEGATIVE VARIATION. (Frog's Gastrocnemius.)

Simultaneous record of a tetanic contraction (white line) and of the accompanying negative variation of a current of injury (black line). (a) The current of injury is normally subsiding; (b) it is suddenly diminished during tetanus (negative variation); (c) it subsequently increases (positive after-variation); and (d) it finally resumes its normal decline. The galvanometer deflection is recorded by a spot of light falling upon a sensitised cylinder revolving round a horizontal axis.

excitation of its nerve, tetanus will include its whole mass, but the change from rest to action will be greater at the uninjured part, A, than at the injured part, B, *i.e.* there will be an action current from B to A, which, while the muscle is tetanised, will diminish the injury current from A to B. This diminution is the '*negative variation.*' A single muscular contraction is sufficient to give a



negative variation, but tetanic contraction will bring it into stronger evidence.

Seeing that a simple negative variation as above described depends upon greater action at an uninjured point, A, than at an injured point, B, it is easy to see that the negative variation will be greater with greater current of injury. The negative variation in muscle appears to precede the contraction to which it belongs, as may be best observed in the heart. (See fig. 187.) We have said 'appears to precede,' because there can be no doubt that under ordinary conditions the mechanical event really begins before it becomes visible, and it is possible that it actually begins with, and not after, the electrical event. In the case of voluntary muscle no interval is demonstrable; both events have a latent period as short as  $\frac{1}{400}$  sec. (B. Sanderson).

*Characters of the negative variation.*—The negative variation in nerve quickly reaches a maximum with increasing strength of stimulating current, is not reversed in direction with reversal in the direction of the stimulating current, does not vary in magnitude with increasing interval between the stimulated and led-off regions of the nerve, and is abolished by ligature in that interval. By the first three of these signs it is distinguished from electrotonic currents; the last is common to both. According to Sanderson and Gotch, the negative variation of potential accompanying a muscular contraction may exceed the potential difference due to injury, *e.g.* 0.08 volt in the former case, as compared with 0.04 in the latter. The negative variation of muscle is greater with greater tension, agreeing thus with the yield of heat and of work, which, as has been stated, are likewise greater with greater tension.

*The double or diphasic variation.*—Whereas the presence of a strong current of injury is favourable to the demonstration of the simple negative variation, which is the sign of the unbalanced negativity of action in the uninjured tissue, it is unfavourable to the demonstration of the diphasic variation, which is a manifestation of the negativity of action, first at one part, then at another part, of uninjured tissue. The double variation depends upon the fact that action is not simultaneous throughout the whole mass of a muscle, but occupies time in its transmission from a point of stimulation.

If A and B (fig. 183) be electrodes applied to an *uninjured* muscle, C the point of application of a stimulus, the part at and near A will commence to act and become negative *before* the part

at and near B; during the interval between the commencement of action at A, and that subsequently at B, the negativity of A is unbalanced; this constitutes the first phase. The part at and near B will continue active and remain negative *after* the part at and near A has ceased to be active; during the interval between the cessation of action at A and that subsequently at B, the negativity of B is unbalanced; this constitutes the second phase. Between the first and second phases, while A and B are both active and negative, the negativity of A more or less accurately balances that at B; this constitutes the isoëlectric interval. Now it is obvious that the slower the transmission of the active negative state from A to B, the more opportunity there is for the manifestation of unbalanced negativity, at A when the process begins, and at B when it ends. Hence it is most easy to demonstrate on the frog's heart, in which the active state travels at the rate of  $\frac{1}{10}$  meter per second, less easy on muscle, in which the rate of transmission is not less than 1 meter, most difficult on nerve, in which the rate of transmission is about 30 meters. The diphasic variation in cooled nerve is, however, easily observed, the rate of transmission being then reduced to about 1 meter per second.

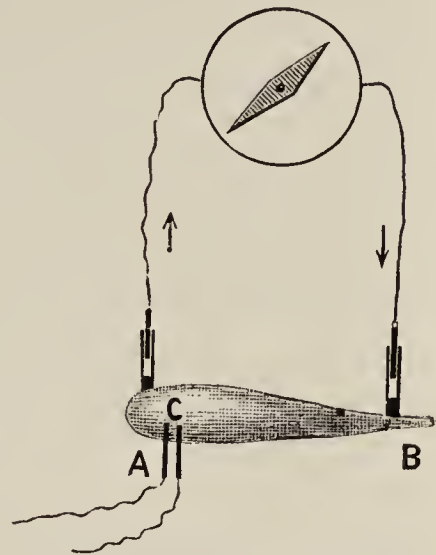


FIG. 183.

We may illustrate the mechanism of a diphasic variation by the following analogy. A railway train (=a nervous impulse) 50 meters long, running at a speed of 50 meters per second, works two signals, A and B, at 10 meters from each other, which fall while the train is at points opposite them, but rise again when the train has passed. During a first period the train is at A, and not yet at B, and A is negative to B (1st or initial phase).



FIG. 184.

During the second period the train is opposite A and B, which are both negative (isoëlectric interval). During a third period the train is past A, but still at B, and B is negative to A (second or terminal phase). With the speed and distances given above



the initial phase would last  $\frac{1}{5}$  sec., the interval  $\frac{4}{5}$  sec., and the terminal phase  $\frac{1}{5}$  sec. Bearing in mind that in a nervous impulse there is no actual transport of matter, but only the propagation of a motion, this figment usefully represents the diphasic and some other of the electrical changes taking place in nerve or muscle—*e.g.* if there is an injury at B, B will be negative to A (current of injury); if a continuous or tetanic series of impulses passes when B is down, A only will fall (excitatory diminution of current of injury); if a continuous series passes when there is no injury, both A and B will be kept down; if impulses pass in an opposite direction, the direction of the phases will be reversed, &c.

The *secondary contraction* (not to be confused with Galvani's second experiment) is caused by a current of action; if, while

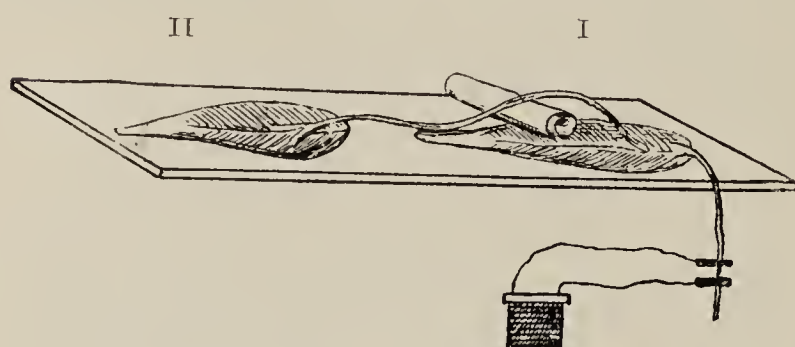


FIG. 185.—THE SECONDARY CONTRACTION.

the nerve of II is resting upon the muscle I, the latter be caused to contract by excitation of its nerve, the sudden electrical variation accompanying the

contraction of I will stimulate the nerve of II and cause a contraction of the muscle II. This is a secondary contraction.

**Experiments on man.**—Two different experiments relating to the electromotive action accompanying muscular contraction have been made upon the human subject—one by du Bois-Reymond relates to voluntary tetanic contraction, the other by Hermann to tetanic contraction caused by electrical stimulation.

*Du Bois-Reymond's experiment.*—The two hands are led off to a galvanometer by two vessels, into each of which a finger is dipped. The voluntary contraction of the muscles of either arm gives a deflection, which indicates the passage of a current through the galvanometer from the passive to the contracting arm, and through the body from the contracting to the passive arm—*i.e.* negativity of the active side. Objection has, however, been taken to the view that the negativity is due to muscular action, and it has been attributed to cutaneous secretory action; it has been found that on a curarised cat, excitation of the sciatic, while causing no muscular contraction, gives the current above described, while on an atropinised cat,

excitation of the sciatic causes muscular contraction, but no action current.

*Hermann's experiment.*—The forearm is led off to a galvanometer by two bracelet electrodes; muscular action is provoked by electrical excitation of the brachial plexus, and the rheotome is employed. A series of observations are taken, the rheotome being shifted from zero (simultaneous excitation and rheotome closure) onwards, which demonstrates the presence of a diphasic variation. The phases are such as to indicate—1, negativity of muscle proximal to the nerve-entrance; 2, negativity of muscle distal from such nerve-entrance. The phenomenon is far more rapid than on frogs' muscle, and time-measurements give for the rate of propagation of the excitatory state a value of about 12 meters per second.

**Experiments on the heart.**—*The frog's heart.* On the frog's heart the normal diphasic variation of its spontaneous systole is as follows:—first phase, base negative to apex; second phase, apex negative to base. If the heart be Stanniused and stimulated at the base, the two phases of the consequent contraction are the same as with the spontaneous beat; but if the stimulus be at the apex, the two phases are reversed, *i.e.* in the first

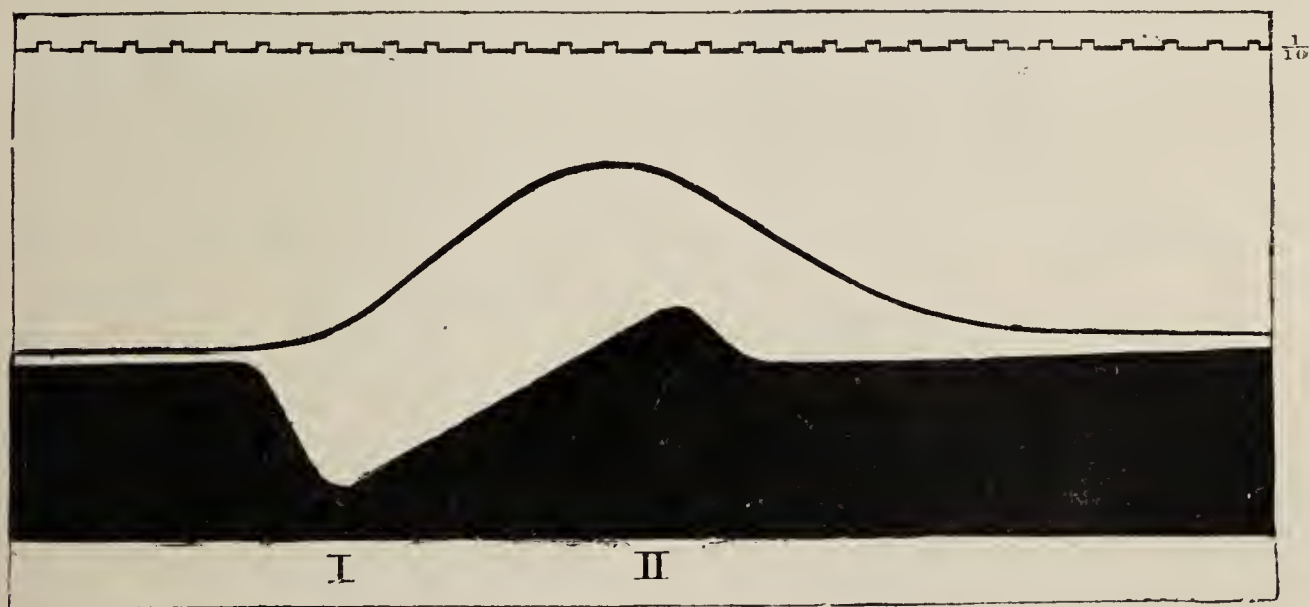


FIG. 186.—FROG'S HEART. DIPHASIC VARIATION.

Simultaneous photogram of a single beat (black line) and of the accompanying electrical change, indicated by the level of the black area, which shows the varying level of mercury in a capillary electrometer. The base of the ventricle is connected with the mercury. I. First phase, base negative to apex. II. Second phase, apex negative to base.

phase apex is negative to base, in the second phase base is negative to apex; in this case the wave of contraction is from apex to base.



The electrical variations of the excised mammalian heart are essentially similar; the normal accompaniment of the spontaneous beat is a diphasic variation in which the two phases are usually such as to indicate—1, negativity of apex; 2, negativity of base. But exceptions to this rule are not uncommon. If, for instance, the apex is injured, the apex phase fails, and the base phase appears alone and increased; if the base is injured, the base phase fails, and the apex phase appears alone; the sequence can be reversed at will by alterations of temperature, and, according to Bayliss and Starling, the sequence of the heart *in situ* is 1, base; 2, apex. By stimulating a normal quiescent ventricle a double variation is produced,



FIG. 187.—CAT'S HEART.  
DIPHASIC VARIATION.

just as in the frog's heart, being  $\begin{cases} 1. A \text{ negative} \\ 2. B \text{ negative} \end{cases}$  if the stimulus is at A, or  $\begin{cases} 1. B \text{ negative} \\ 2. A \text{ negative} \end{cases}$  if the stimulus is applied at B.

*The human heart (w.).*—The electrical variations of the

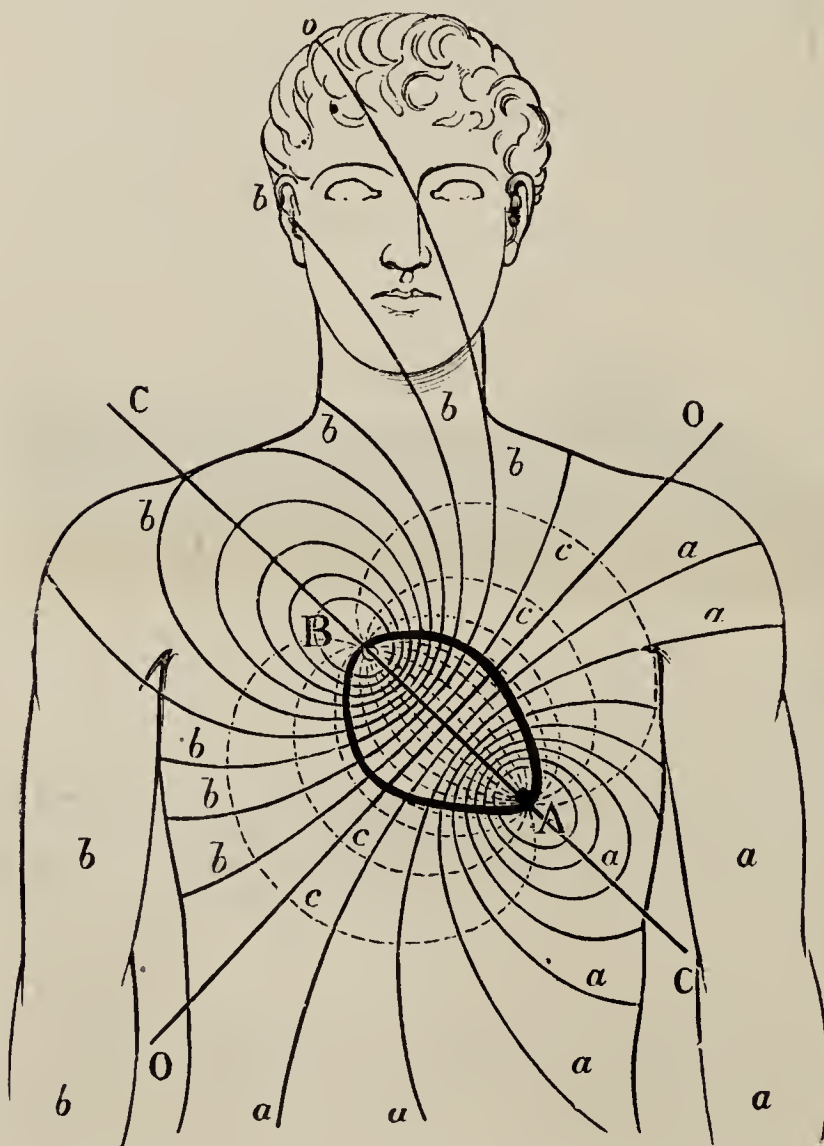


FIG. 188.

Let A and B respectively represent apex and base of the ventricular mass. Then, if at any moment a difference of potential should arise between A and B, a current, *c c c*, will be established along and around the axis *C C*; the line *O O* will represent the plane of zero potential or equator; the lines *a a a*, *b b b* will represent equipotential curves around A and B. A difference of potential between A and B will be manifested if two leading-off electrodes are applied on opposite sides of the equator, *O O*; no such difference will be manifested if both electrodes are on the same side of the equator. The equator *O O* will divide the body into two asymmetrical parts, (1) a portion *b b b* including the head and right upper extremity, (2) a portion *a a a* including the three other extremities.

mammalian heart may also be studied upon intact animals

and upon man by means of Lippmann's capillary electrometer (p. 314). By means of this instrument variations of electrical potential that are associated with the beat of the human heart can be demonstrated, and their mode of distribution in the body determined, by simply 'leading off' from various points of the surface. This mode of distribution will best be understood by reference to the accompanying diagram and description.

If we lead off to the electrometer by any two points, *a* and *b*, on opposite sides of the equator (*e.g.* by the mouth and the left hand), we shall see the mercury pulsate with each beat of the heart; if we lead off to the electrometer by any two points, *a a* or *b b*, on the same side of the equator (*e.g.* by the mouth and the right hand), the electrometer will not be affected by the heart-beat.

On careful inspection of the electrometer, or better, by taking photographic records, it will be found that each beat of the heart is attended with a double movement of the column of mercury. The physiological significance of this double movement is, however, difficult to determine; usually, and with a good electrometer, the two movements are apparently in the same direction; the direction of the second movement, which is comparatively slow, is such as to indicate negativity of *b*; that of the first movement,

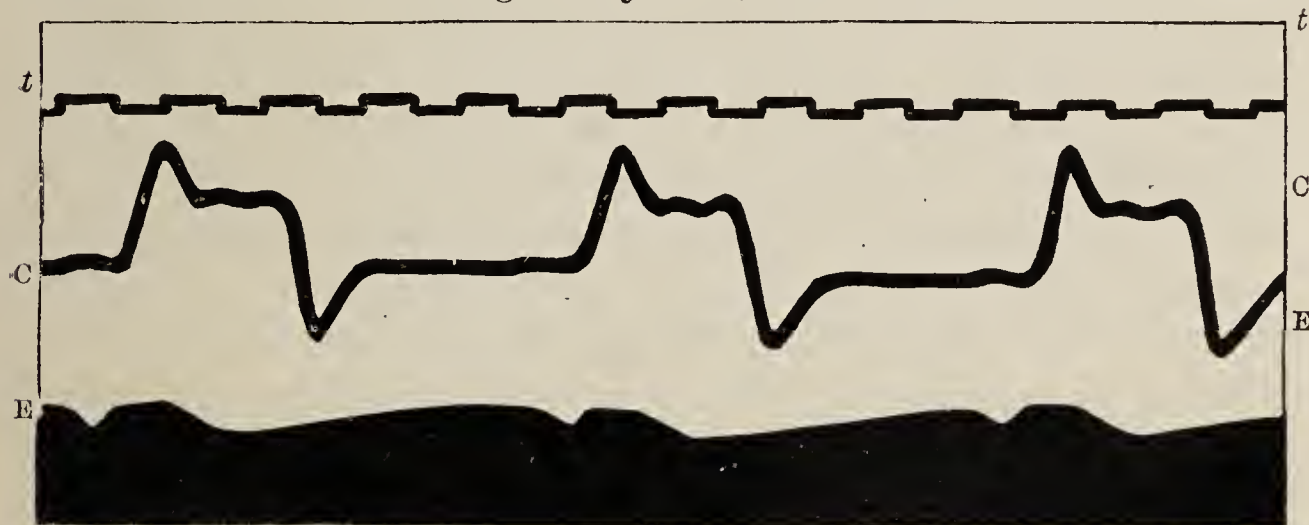


FIG. 189.—HUMAN HEART. DIPHASIC VARIATION, E E; AND SIMULTANEOUS CARDIOGRAM, C C.

The leads-off to the capillary electrometer were from the mouth to the sulphuric acid, and from the left foot to the mercury. Time, *t t*, marked in  $\frac{1}{10}$ ths sec.

which is comparatively rapid, is not easy to determine; it is probably itself a double movement due to negativity of *b* suddenly replaced by negativity of *a*. Thus the order of negativity is apparently *b.a.b.*, and not *b.a.* or *a.b.* A similar triphasic variation, *b—a—b*, in place of the ordinary variation, *b—a*, is often to be observed in the spontaneous beat of the frog's heart, the final *b* phase being presumably due to the contraction of the bulbus arteriosus. It is therefore probable that in all kinds of



hearts the contraction begins from the base, spreads to the apex, and ends at the base.

*Secondary contraction from the heart* is obtained when the heart is substituted for the muscle I in fig. 185. Just before the commencement of each beat the nerve of II is excited by the electrical variation that belongs to the contraction of the heart, and the muscle II contracts. Under favourable circumstances (*i.e.* with a very excitable nerve and an uninjured heart giving a diphasic variation) the muscle contracts at the end as well as at the beginning of the heart's contraction. Occasionally, on decapitated animals, immediately after decapitation, the left side of the diaphragm may be observed to contract at each beat of the heart. This is due to excitation of the left phrenic nerve by the electrical variation with cardiac contraction, and is an instance of secondary contraction from the heart.

Kühne has shown that if two muscles are pressed together, excitation of either will cause contraction of both; this is secondary contraction from muscle to muscle. Secondary contraction from nerve to nerve has been demonstrated by Hering on 'cooled frogs.'

**Cutaneous currents, glandular currents.**—The skin of all animals, not excepting fishes, is in the normal state traversed by an electrical current from without inwards, mainly caused by cutaneous glands. According to Hermann, the glandless skin of the eel gives a current of  $\cdot 003$  to  $\cdot 007$  Dan., attributable to the degenerative action or partial death that accompanies the formation of the epidermis. Such skin has no distinct glands, but it contains large numbers of single secreting cells, and cannot, therefore, be considered as a non-glandular tissue. According to Reid, the normal current from outer to inner surface is increased by excitation, *i.e.* it gives a positive variation, and this excitatory increase of current is temporarily reduced or abolished by chloroform, by carbon dioxide, and by atropin. According to Biedermann, the variation is positive when the skin is cooled or dried, and negative if it is warm and moist.

The best-known instance of a cutaneous gland current under the influence of secreto-motor nerves is that which is afforded by the pad of the cat's foot, where excitation of the sciatic nerve causes an outbreak of sweat accompanied by a positive variation of the incoming skin current, effects which are no longer produced after the administration of atropin. The current obtained in du Bois-Reymond's experiment on man is, according to

Hermann, of a similar nature, *i.e.* a gland current directed from without inwards in the skin of the active hand. According to Tarchanoff, the cutaneous currents of man are exceedingly sensitive, especially in parts of the skin where the sweat-glands abound, being increased by all kinds of cerebral activity in the absence of visible muscular contractions.

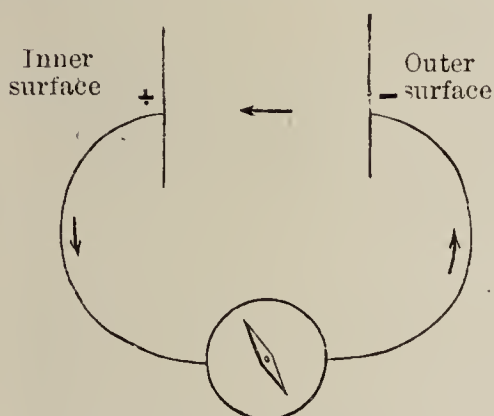


FIG. 190.—SKIN CURRENT.

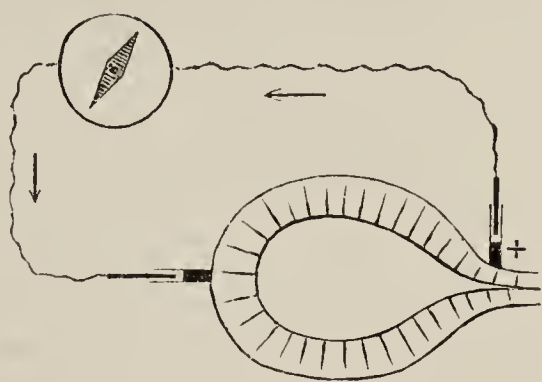


FIG. 191.—GLAND CURRENT.

The submaxillary gland is also the seat of a glandular current which, according to Bayliss and Bradford, is directed from without inwards, or in the galvanometer circuit from hilum to surface, and which is increased by excitation of the chorda tympani (positive variation), diminished by excitation of the sympathetic (negative variation). Bohlen finds that excitation of the vagus causes an electrical change of the stomach-wall, attributable to secretory action.

**Cerebro-spinal currents.**—Caton, in 1875, published observations to show that visual excitation causes electrical effects in the cortex cerebri. According to Setschenow, the medulla oblongata of the frog, if led off to the galvanometer, exhibits irregular series of electrical variations, which may be excited or arrested by excitation of the central end of the sciatic. Gotch and Horsley have recently led off the electrical signs of nerve-action from the white columns of the spinal cord by means of the capillary electrometer. More recently still, Beck and Cybulsky have made observations similar to those of Caton; and v. Fleischl has published observations relating to the cerebral currents on man. All that we may safely conclude from these several observations is that the current of action occurs in the *white* matter of the cord, as in an ordinary nerve; its manifestation by the *grey* matter of nerve-centres is still an open question.

**Retinal currents.**—If two unpolarisable electrodes connected with a galvanometer are brought in contact with an excised frog's eye, one with the cornea, the other with the fundus, a deflection



is obtained showing that the fundus is negative to the cornea. This is an ordinary current of injury arising from the cut optic nerve, and not assignable to the retina. If an isolated eyeball, or better, an isolated retina, is put in connection with the gal-

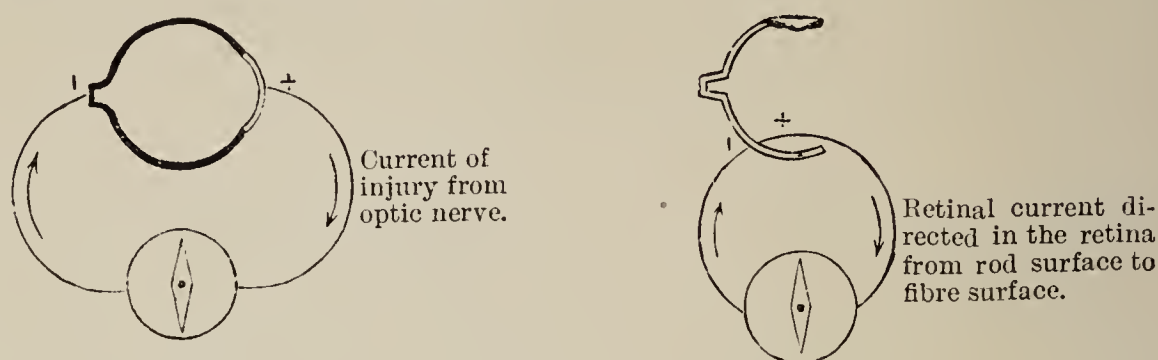


FIG. 192.—EYE CURRENTS.

vanometer, and kept in the dark, then illuminated for a short period of time (15 to 30 secs.) by opening a shutter, the current will be found to undergo a variation—(1) at the commencement, (2) during the incidence, (3) at the termination of the illumination, as represented in fig. 193.

An entirely satisfactory explanation of these effects cannot be given, but the important fact remains that light acting upon the retina causes an electrical change which is the physical token of retinal activity. Holmgren showed by exclusion that the electrical variation caused by light depends upon the retina only, and is not due to the action of light upon any other constituent of the eyeball; nor is it due to changes of pigment. He showed

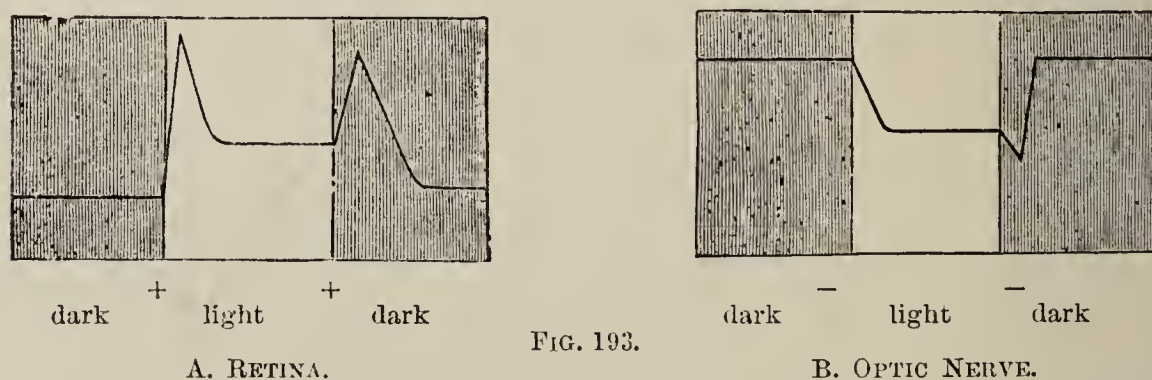


FIG. 193.

A. RETINA.

B. OPTIC NERVE.

The rod surface is negative to the fibre surface; with illumination this negativity is increased at first, then less increased (in some cases actually diminished); at cessation of light a second increase occurs. (Kühne and Steiner.)

The transverse section is negative to the longitudinal surface; with illumination this negativity is diminished; at cessation of light it undergoes a second diminution, and then returns to its original magnitude. (Kühne and Steiner.)

also that the external or choroidal surface of the retina is negative to its internal surface. Kühne and Steiner found that the variation occurs nearly as well on a retina devoid of visual purple as on a non-bleached retina, although its character was somewhat different in the two cases. Dewar and McKendrick found that

white or coloured light acted upon the retina in the following order of strength—white, yellow, green, red, blue. They also demonstrated the electrical variation on intact animals and on man, one electrode being on the eye, the other on any other part of the body. With regard to the magnitude of the change, they estimated it at  $\frac{1}{10000}$  Daniell. The essential phenomenon is the current of action; whether or no a current pre-exists, due to injury or other accidental conditions, is a matter of secondary importance.

**Electric fishes.**—The most striking manifestations of animal electricity are those that are afforded by electric fishes—*gymnotus*, *torpedo*, and *malapterurus*. The electric organs are practically powerful batteries, for which a succession of instantaneous discharges of high intensity can be liberated at will of the animal, or in a reflex manner by cutaneous stimulation, or experimentally by direct electrical excitation of the organ itself or of its efferent nerves. An electrical organ is in effect a terminal apparatus of nerve, analogous with muscle, giving action currents of very great intensity, and, like muscle, alkaline in the normal

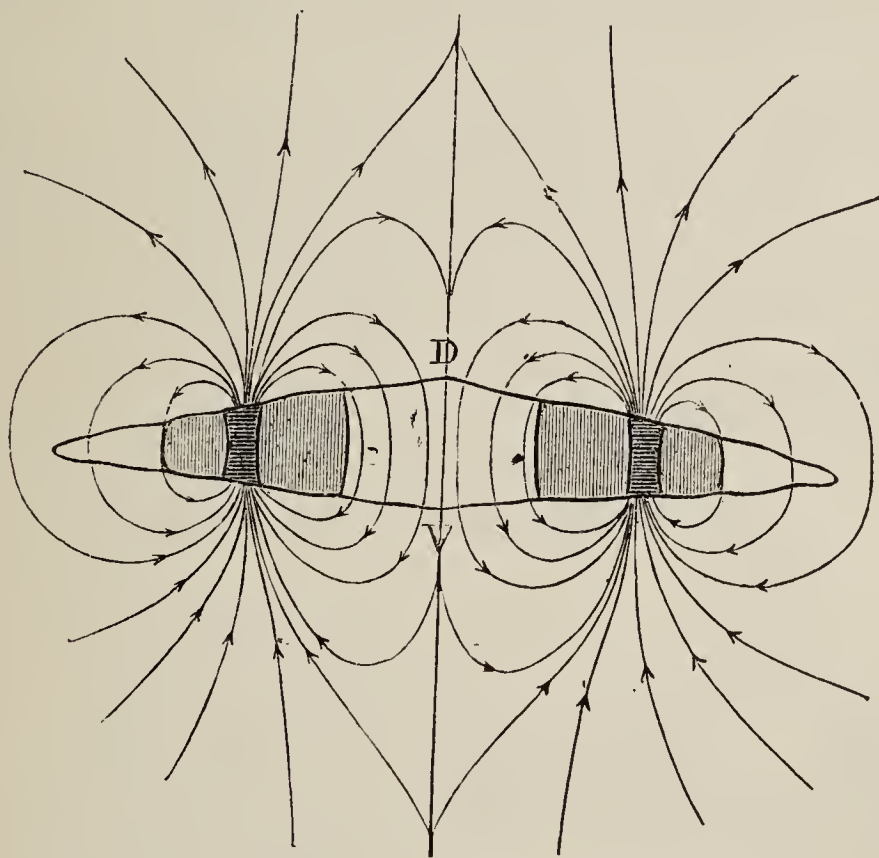


FIG. 194.—DIAGRAMMATIC TRANSVERSE SECTION OF TORPEDO.

D dorsal, V ventral surface; Direction of discharge indicated by arrows.  
(Du Bois-Reymond.)

state, acid after death or excessive activity. In the torpedo it is formed of two lateral masses composed of numerous hexagonal prisms perpendicular to the surface of the body, these prisms



being subdivided into compartments by transverse septa parallel with the surface. In the discharge of the organ the current flows from the dorsal to the ventral surface through the galvanometer, from the ventral to the dorsal surface through the organ itself. According to du Bois-Reymond, the electrical resistance

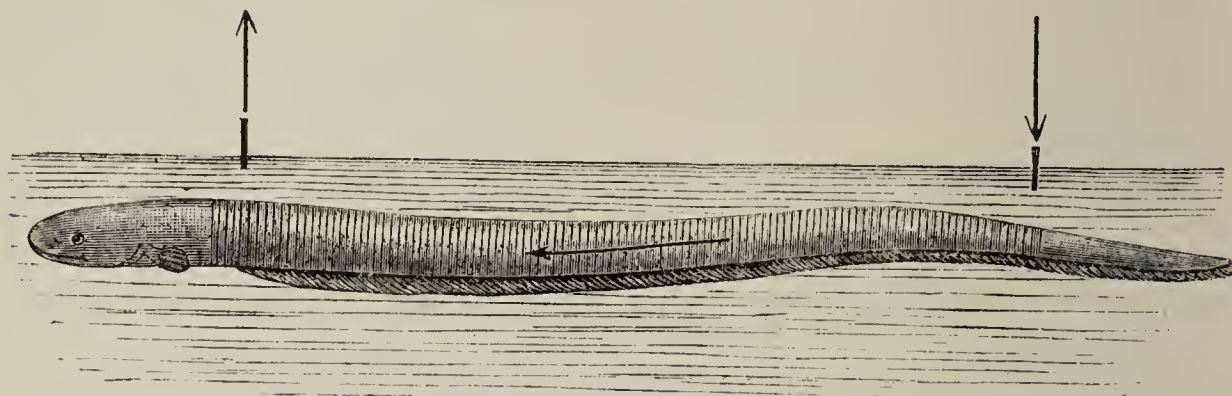


FIG. 195.—GYMNOTUS.

Direction of discharge indicated by arrows and leading-off electrodes from water.

of the living organ is very much greater to *down* currents from dorsal to ventral surface than to *up* currents from ventral to dorsal surface. According to Gotch, this apparent difference is the effect of an upward action current adding to the up current and subtracting from the down current.

In the gymnotus the shocks are always directed from tail to head in the animal, in the malapterurus they are directed from head to tail. The direction, as was pointed out by Pacini, depends upon the points of entrance of the electric nerves; in the torpedo they are distributed to the ventral surface, in the gymnotus to the posterior surface, and in the malapterurus to the anterior surface of the organ. The discharge is always such that the

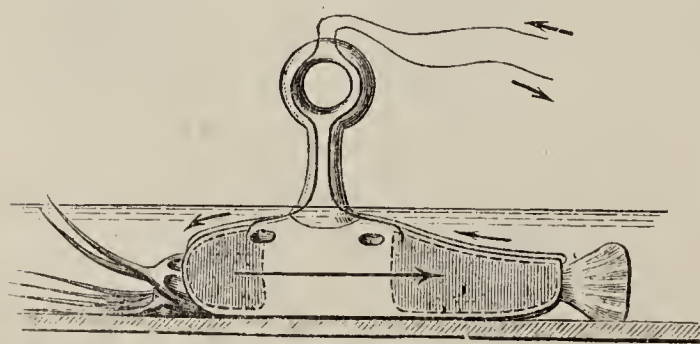


FIG. 196.—MALAPTERURUS.

Led off by two shields; discharge indicated by arrows. (Du Bois-Reymond.)

nerve-surfaces are active, *i.e.* analogous with the zinc of a Daniell cell. The discharge is a discontinuous one, composed of as many as 200 shocks per second in the fresh condition, but with exhaustion falling in frequency and in force. A single shock has a latent

period of about  $\frac{1}{200}$  second, and a duration of  $\frac{7}{100}$  second. The rate of transmission along the electrical nerves is about 7 meters per second; the fish themselves are unaffected by electrical currents sufficient to kill non-electrical fish.

*Physiological modifications are conveniently gauged by their electrical signs* (w.).—The current of action in an excitable tissue is a sign of action, and the magnitude of such current aroused by excitation may, with certain reservations, be taken as an index to the magnitude of action; the value of such indications is greatly enhanced if they are photographed in a continuous record. The tissues most suitable to the study of the relations between excitation and activity under various conditions are muscle, nerve, and retina. The most significant experiment on *muscle* consists in a verification of the parallelism between its mechanical action and the accompanying electrical changes. Any simultaneous record of the mechanical and electrical effects in voluntary or in cardiac muscle exhibits this correspondence; a more or less strongly contracting muscle gives a more or less strong negative variation, cardiac muscle made to relax by vagus excitation gives a positive variation; muscular fatigue is as well (or better) recorded by a galvanograph as by a myograph.

As regards the *retina*, in addition to the fundamental fact that an electrical change is provoked by light, two points are most worthy of note.

1. Equal increments of stimulation cause diminishing increments of electrical effect, so that the cause-effect curve rises concave towards its abscissa like the Weber-Fechner curve (p. 555).

2. Prolonged series of electrical effects, caused by strong illumination at regular intervals, exhibit no signs of fatigue, unlike muscle in this respect, but like nerve.

Experiments on **nerve** are in many respects the most convenient and broadly instructive; electrical signs of activity are the only available index of action within nerve itself, apart from its terminations, and isolated nerve is practically the most convenient representative of living, *i.e.* excitable, matter

in general. To study the effect of reagents (anæsthetics, narcotics, sedatives, neutral salts, alkaloids, &c.) upon the current of action, is to study their effect upon the 'excitability' of living

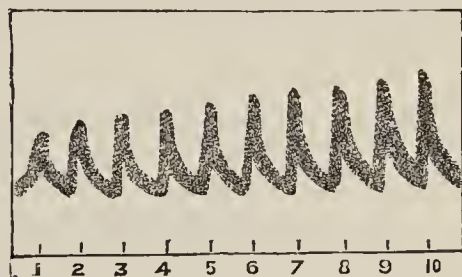


FIG. 197.

Galvanometric deflections (*i.e.* retinal effect) in a frog's eyeball caused by exposures, each lasting 7.5 secs. to the light of  $\frac{1}{100}$ ,  $\frac{2}{100}$  to  $\frac{10}{100}$  of a standard candle. The curve of increasing magnitude of deflection with arithmetically increasing stimulus closely resembles the Weber-Fechner curve of fig. 300.



matter. The chief facts that can be briefly stated in this connection are:—

1. Prolonged series of action-currents caused by tetanisation at regular intervals, exhibit no diminution, *i.e.* nerve is practically inexhaustible.

2. Within the physiological working range of nerve (*i.e.* within the range between minimal and maximal response of its muscle), the negative variation increases arithmetically with arithmetically increased strength of stimulation. The line of increase, unlike that of muscle and of the retina, is a straight line.

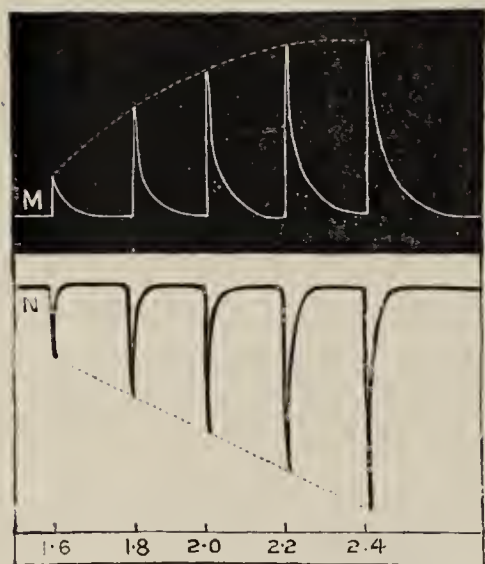


FIG. 198.

Diagram of muscular contractions M, and of simultaneous negative variations of nerve N (central end), with stimuli increasing from 1.6 to 2.4 units.

3. Anæsthetics (ether and chloroform) temporarily suppress the action currents, *i.e.* temporarily abolish excitability. Suppression by chloroform is more prolonged than suppression by ether, and is frequently final, *i.e.* not followed by recovery. The recovery after ether is frequently followed by secondary augmentation.

4. Carbon dioxide in small quantity, *e.g.* 4 per cent. as in expired air, augments the action-currents, *i.e.* increases excitability, in large quantity causes primary diminution or abolition followed by secondary augmentation, *i.e.* acts like ether.

5. Prolonged tetanisation gives rise to augmentation of the action currents, *i.e.* produces an effect similar to that brought about by a small quantity of CO<sub>2</sub>.

6. Oxygen, nitrogen, hydrogen, nitrous oxide, carbon monoxide, have no marked effect upon the action currents.

From these and other data<sup>1</sup> we may draw two conclusions of principal importance:—

1. That the tetanisation of nerve is attended with an evolution of carbon dioxide.

2. That the inexhaustibility of nerve (and of the retina) is due to active reintegration rather than to sluggish disintegration.

<sup>1</sup> Cf. *Phil. Trans. R. S.* (Croonian Lecture), 1896, and other papers mentioned in the Bibliography

A negative variation during excitation is frequently suc-

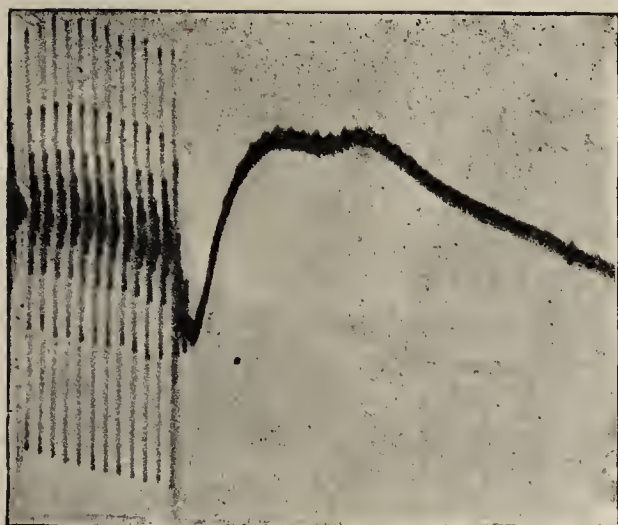


FIG. 199.—ACTION OF CHLOROFORM.

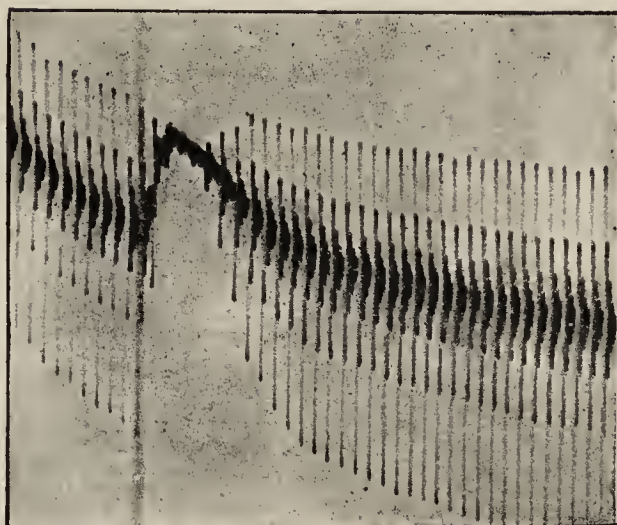
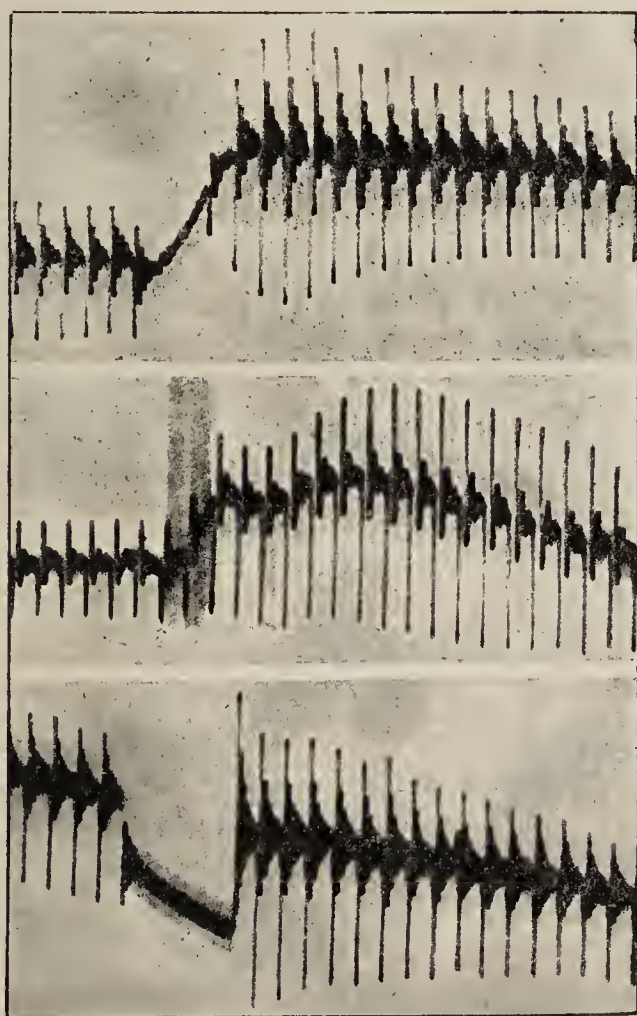


FIG. 200.—ACTION OF ETHER.



Full effect of carbon dioxide.  
(Abolition followed by  
augmentation of the electrical  
response.)

Slight effect of carbon dioxide.  
(Primary augmentation of the  
electrical response.)

Effect of 5 min. tetanisation.  
(Primary augmentation of the  
electrical response.)

FIG. 201.

Records of the action of some reagents upon the negative variation of nerve. The response is temporarily suppressed by ether, permanently abolished by chloroform, temporarily suppressed by 'much' carbon dioxide (with subsequent augmentation), augmented by 'little' carbon dioxide, augmented by prolonged tetanisation. The nerve is enclosed in a moist chamber, excited at 1 min. intervals through one pair of electrodes, 'led off' to the recording galvanometer by a second pair of electrodes. The time during which the ether or chloroform vapour or carbon dioxide is blown into the nerve-chamber is in each case indicated by the dark vertical bar. (w.)

ceeded by a positive variation after excitation has ceased: the former being probably significant of disintegration, the latter



possibly significant of reintegration; and under certain conditions a positive variation may be witnessed during excitation. This positive variation, which it is difficult to distinguish from an anelectrotonic effect, is subject to the action of anæsthetics; it is diminished or reversed under the influence of  $\text{CO}_2$ , and in consequence of prolonged tetanisation.

The effect of reagents in solution is no less easily ascertained than that of gases. The nerve is put to the question before and after immersion in the solution, and the alterations of its response give measure of the influence of the drug. We may in this way determine whether a given chemical molecule acts or no upon the nerve-protoplasm, and—in the affirmative case—what portion of the molecule is more or less influential. *E.g.* it having been found that potassium bromide,  $\text{KBr}$ , is markedly depressant, the question whether the basic moiety  $\text{K}$  or the acid moiety  $\text{Br}$  plays the greater part, will be answered by experiments with  $\text{NaBr}$  and  $\text{KCl}$ . The former is ineffective, the latter is effective, from which it must be concluded that in the depressant action of  $\text{KBr}$ , the basic element  $\text{K}$  is the predominant partner. Among the alkaloids, there may be named in order of efficacy: aconitine and veratrine as very effective, curarine and digitaline as effective, atropine and aconine as ineffective.

## CHAPTER XII

### LIGHT AND VISION

#### PHYSICAL DATA

#### LIGHT

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*Reflection.*—A ray of light falling upon a polished surface is *reflected* (fig. 202). B N being the normal or perpendicular to the reflecting surface, S S, at the point of incidence, B, the angle of incidence, A B N, is equal to the angle of reflection, N B C.

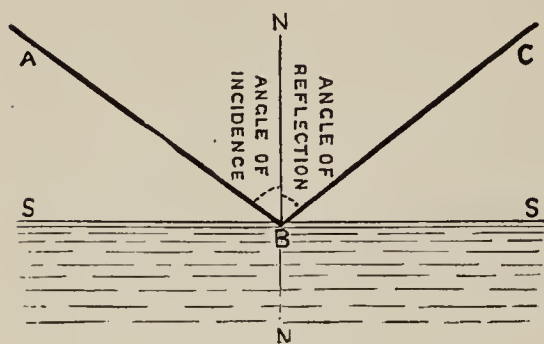


FIG. 202.—REFLECTION.

*Refraction.*—A ray of light passing from a medium of less density into one of greater density, or *vice versa*, is bent or *refracted* from its original direction (except its direction be perpendicular to the surface of contact between the two media). Passing from a medium of less to one of greater density, *e.g.* from air to glass, the ray is refracted towards the normal; passing from a medium of greater to one of less density the ray is refracted away from the normal.

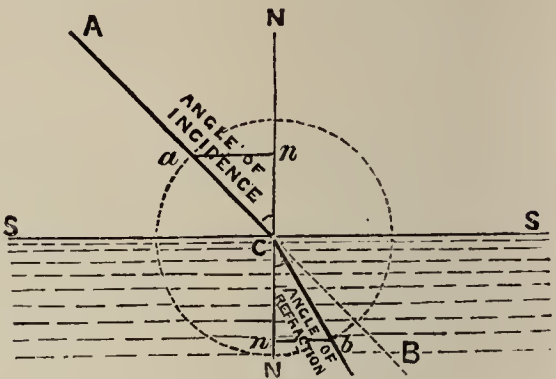


FIG. 203.—REFRACTION.

S S, boundary surface between air and glass. N N, perpendicular to the surface S S. A ray of light passing in the direction A C B is deflected towards N N in passing from air to glass. A ray of light passing in the direction B C A is deflected from N N in passing from glass to air. A C is the incident ray. C B is the refracted ray.

The relation  $\frac{\sin \text{angle of incidence}}{\sin \text{angle of refraction}}$  or  $\frac{a n}{b n}$  (fig. 203) is the *index of refraction* of the substance into which the ray passes from air. It is constant for all angles of incidence, but various for various refracting media. Of glass, the index of refraction is 1.5 to 1.6, according to the kind of glass.

Of water . . . . .	1.33
„ crystalline lens . . . . .	1.40 to 1.45
„ vitreous body . . . . .	1.34
„ aqueous humour . . . . .	1.34

If the ray pass through a glass plate with parallel surfaces (fig. 204), it is bent towards the perpendicular at B, from the perpendicular at C. Its direction from C to D is parallel with its direction from A to B.

If the ray pass through a glass prism (fig. 205), it is refracted at B towards the perpendicular N, and at C from the perpendicular N—*i.e.* in each case towards the base of the prism.

A pair of prisms joined by their bases will deflect rays, in the same plane, passing through each of the two prisms towards each other. A pair of prisms joined by their apices will deflect rays passing through each of the two prisms away from each other.

**Lenses.**—A biconvex lens may be regarded as composed of an infinite number of prisms with their bases towards the centre of the

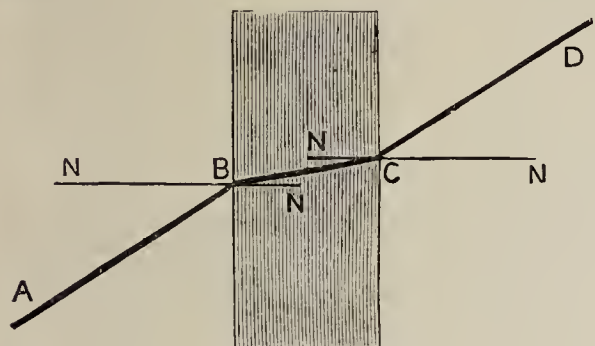


FIG. 204.—RAY THROUGH PLATE.

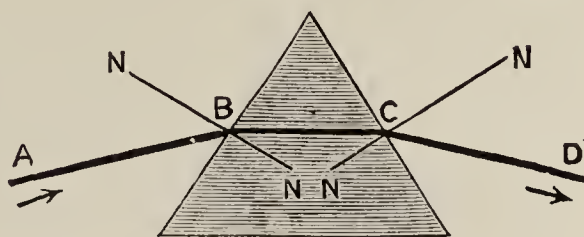


FIG. 205.—RAY THROUGH PRISM.

lens; its effect will be to cause rays of light to converge. A biconcave lens may be regarded as composed of an infinite number of prisms with their apices towards the centre of the lens; its effect will be to cause rays of light to diverge. The *focus* of a lens is the point of

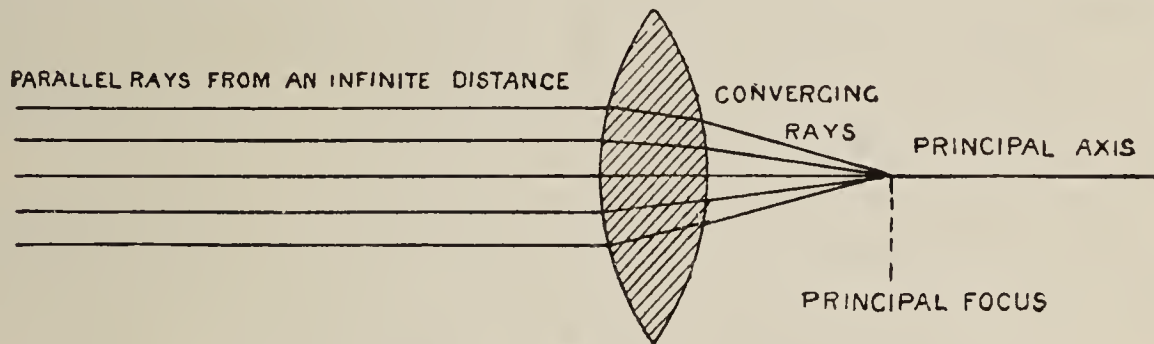


FIG. 206.

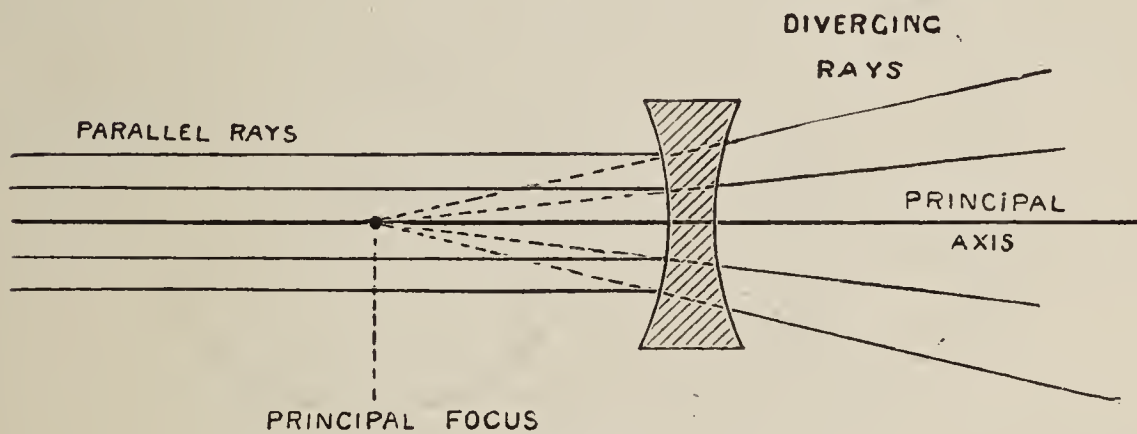


FIG. 207.

meeting of refracted rays (real focus), or of their prolongation backwards (virtual focus). The *principal focus* is the focus of rays parallel with the principal axis of the lens, and its distance from the lens is the principal focal distance.

**Conjugate foci.**—A conjugate focus is the focus of divergent rays from a luminous point.  $c$  and  $c_1$  (fig. 208) are conjugate points. The principal and conjugate foci are *real* foci. A *virtual* focus is formed by a luminous point situated between the lens and its principal focus. The rays after passing through the lens are still divergent; their prolongation backwards to their point of intersection gives the virtual focus of the point. A concave lens causes divergence of the rays of light by which it is traversed, and has only virtual foci (fig. 207).



*Centre of curvature.*—A spherical surface—*i.e.* a surface forming part of a sphere—has a centre of curvature,  $P$ , which is the centre of the sphere, and a radius of curvature, which is the distance between centre

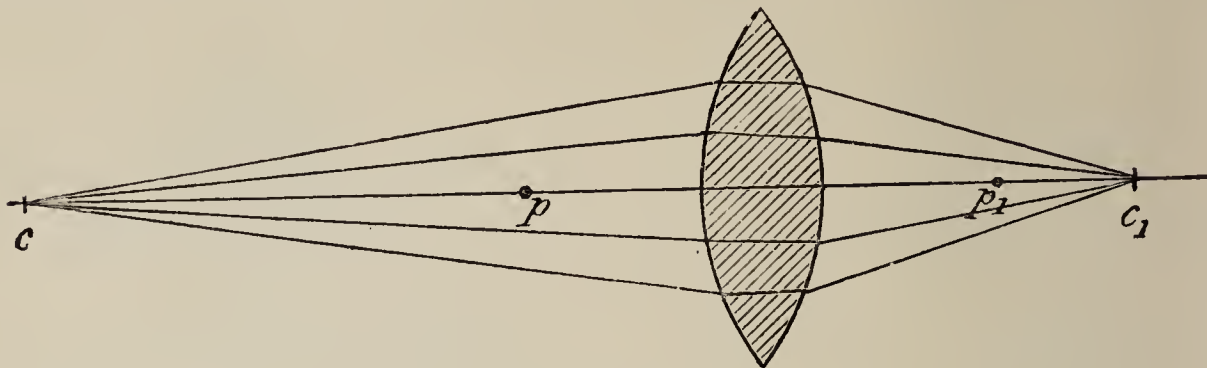


FIG. 208.

and surface. A straight line joining the centres of curvature of the two spherical surfaces of a biconvex lens is its *principal axis* (fig. 209). The point of the principal axis through which any luminous ray,  $SS$ , passes without angular deviation, the emergent being parallel to the incident ray, is called the *optical centre* of the lens; any straight line,  $SS$ , passing through the optical centre, without passing through the centres of curvature, is a *secondary axis*.

Rays parallel with a secondary axis, or emitted from points situated on a secondary axis, form foci, just as do rays parallel with the principal axis; hence images can be formed.

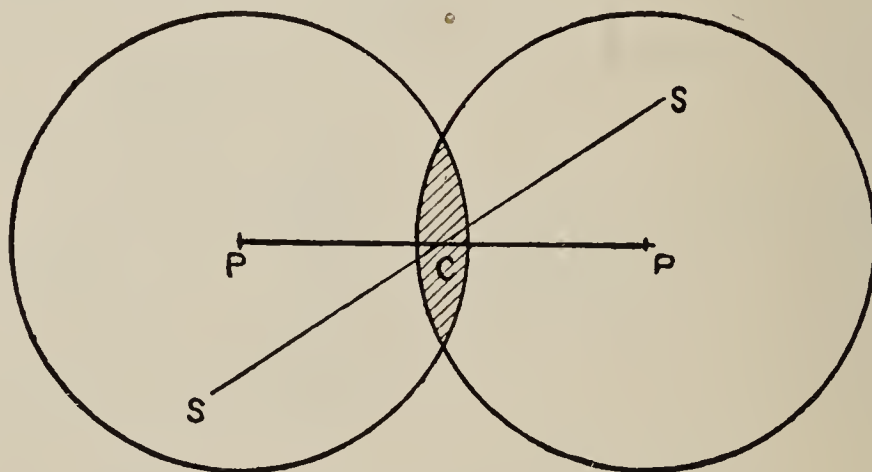


FIG. 209.—CONSTRUCTION OF A BICONVEX LENS.

The points  $PP$  are its centres of curvature. A line joining these points is the principal axis.  $C$ , the middle point of this line, is the centre of the lens; any line,  $SS$ , passing through  $C$  is a secondary axis.

**The image** of an object is the collection of the foci of its several points. A collection of real foci constitutes a *real image*; a collection of virtual foci constitutes a *virtual image*.

If an object be situated beyond the principal focus of a lens, it forms an image which is real and inverted. If it be far from the principal focus, the image is small and close to the principal focus. If it be near the principal focus, the image is large and distant from the lens. These two extreme cases are respectively illustrated by a camera or an eye, and by a magic-lantern. If the object be at the principal focus, no image is formed. If it be between the lens and its principal focus,

there is no real image formed, but a virtual image, erect and larger than the object. The nearer the object is to the principal focus, the larger and more distant its image.<sup>1</sup> A real image may be projected on a screen (large), or received by the eye (small); a virtual image can only be seen when the prolongations of its constituent rays fall on the eye.

*Mirror images.*—The rays reflected from convex mirrors are divergent; such mirrors can therefore have only virtual foci, and can only give a *virtual image*, which is *erect and smaller* than the object; the greater the convexity, the smaller the image. Concave mirrors give a *virtual erect image* of an object which is situated between the mirror and its principal focus; a *real reversed image* of an object situated farther from the mirror than its principal focus. If the object is between the principal focus, F, and centre of curvature, C, its image is larger than the object, and situated beyond the centre of curvature; and, *vice versa*, if the object is situated beyond the centre of curvature, its image is smaller than the object, and situated between the principal focus and centre of curvature; the image and the object occupy 'conjugate' positions.

Sanson's images are mirror images from the cornea and lens. The first and second images are virtual erect images reflected by convex surfaces; the third is a real reversed image reflected by a concave surface (p. 429).

*Spherical aberration.*—A spherical lens does not bring to an exact focus all the rays passing through it; rays passing through the central part of the lens are less strongly refracted than rays passing

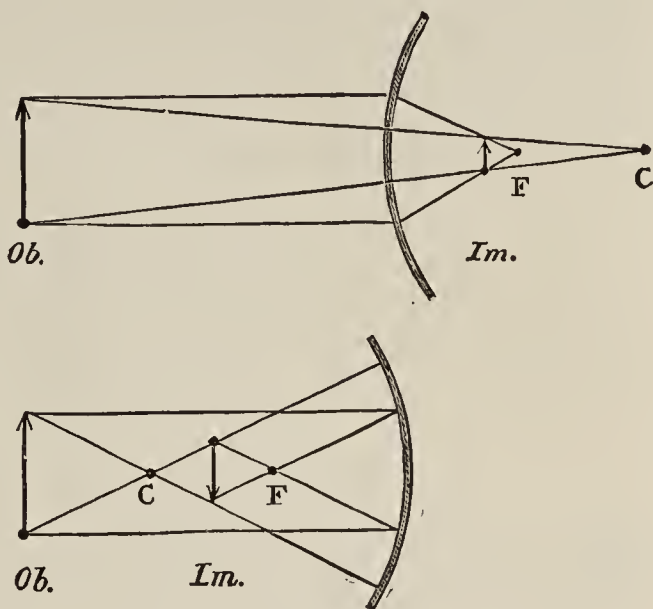


FIG. 210.

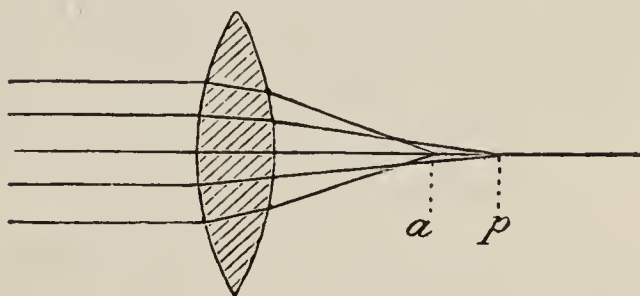


FIG. 211.—SPHERICAL ABERRATION.

Rays through central part of lens focussed at *p*, rays through border of lens focussed at *a*.

<sup>1</sup> The position of conjugate foci (or images) is calculated from the formulæ—

$$\frac{1}{d} + \frac{1}{d'} = \frac{1}{f} \text{ for real focus of convex lens or concave mirror.}$$

$$\frac{1}{d} - \frac{1}{d'} = \frac{1}{f} \text{ for virtual focus of " " "}$$

$$\frac{1}{d'} - \frac{1}{d} = \frac{1}{f} \text{ for virtual focus of concave lens or convex mirror.}$$

The size of the image is to the size of the object as the distance *d* of the object is to the distance *d'* of the image.



through parts farther from the centre. This interferes with the distinctness of images, and is in part corrected by a diaphragm—*i.e.* a screen with a central aperture so that only central rays traverse the lens, and peripheral rays are arrested. In the crystalline lens of the eye this is further corrected by a peculiarity of optical property—the refractive index is greatest at the centre.

*Chromatic aberration.*—A lens does not refract the constituents of white light in precisely equal degrees; it refracts rays of short wave-length more strongly than rays of greater wave-length—*e.g.* the violet more strongly than the red rays, so that the focus of violet rays is closer to the lens than that of red rays. Hence the image of a spot of

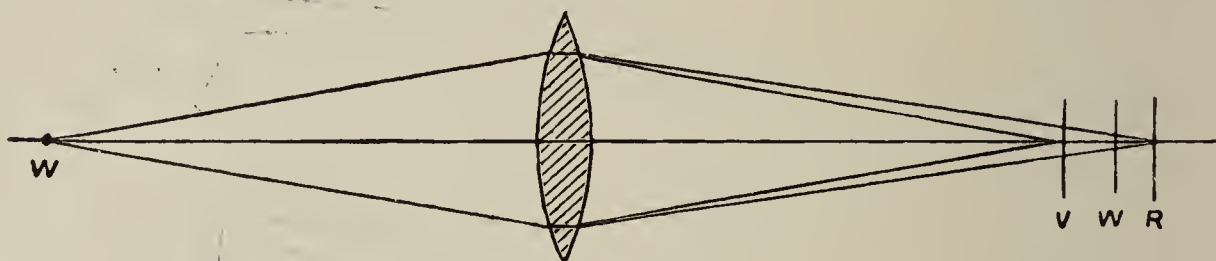


FIG. 212.—CHROMATIC ABERRATION.

light is apt to be surrounded by a red halo when the screen is rather within the mean focal distance of the lens, by a violet halo when the screen is rather beyond the mean focal distance of the lens. Chromatic aberration is an instance of the dispersion of white light by a prism into the colours of the spectrum.

*Lens units. The diopter.*—The stronger a lens, the shorter its focal distance, and *vice versa*. The strength of a lens may therefore be indicated by stating its focal length—*e.g.* lenses of 1, 2, 3, &c. inches focal length are progressively weaker, their strengths being in the proportion 1,  $\frac{1}{2}$ ,  $\frac{1}{3}$ , &c. This is, however, an inconvenient scale in actual practice, as, for instance, in the numeration of spectacle lenses, or in the estimation of the refraction of the eye.

It is more convenient to adopt as the unit a weak lens, viz. a lens with a focal distance of 1 meter. This unit is called the *diopter*, and is written 1 D. A lens twice as strong has a converging power equal to two such lenses, its strength is expressed as 2 D, and its focal distance will be  $\frac{1}{2}$  meter. A lens ten times as strong has a converging power of 10 D, its focal distance is  $\frac{1}{10}$  meter, *i.e.* 10 cm.

The focal value of concave lenses is similarly expressed with the negative sign prefixed; for the value of convex lenses the positive sign is commonly prefixed; thus, +  $n$  D is the power of a convex lens, —  $n$  D is the power of a concave lens.

COLOUR<sup>1</sup>

White light is composed of many kinds of light—red, orange, yellow, green, blue, violet—and intermediate between these named colours innumerable others. This fact is demonstrated by interposing a prism on the path of a pencil of sunlight. Refraction occurs, being greatest for violet rays, least for red rays; the white light is decomposed into its constituent colours, and a continuous spectrum of the prismatic colours is obtained.

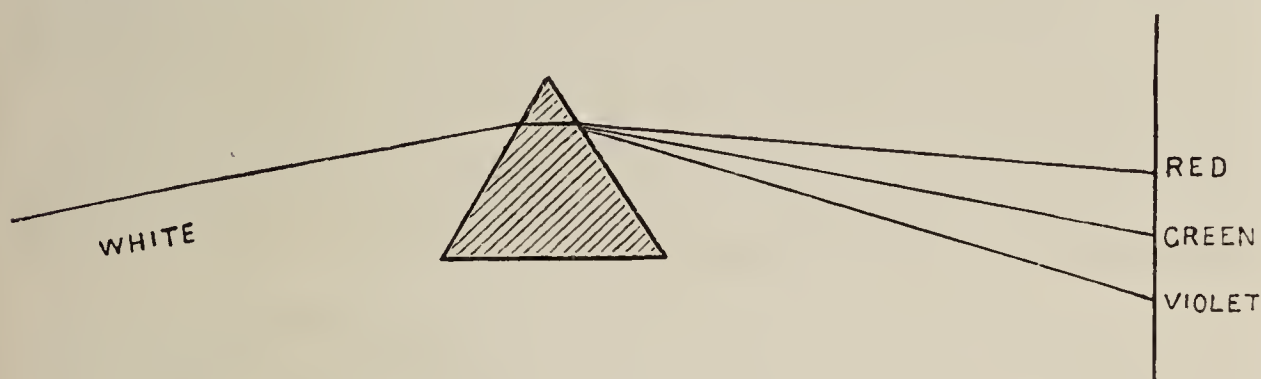


FIG. 213.

White light may be recomposed from its constituents, either *physically*, as by a second prism reversing the dispersion produced by the first prism, or *physiologically*, by causing the prismatic colours to fall upon the retina in sufficiently rapid succession. The readiest method of studying the effects of different colour-mixtures is by means of a revolving disc to which two or more coloured discs are fixed, overlapping more or less, so that varying proportions of the colours occupy the entire circumference of the disc (Maxwell).

White light is composed of an infinite number of simple colours; it can, however, be produced by mixture of the three primary colours, or even of only two colours in certain proportions, but in this case one at least of the two colours must be other than a primary. Any two colours which, taken together in proper proportions, form white, are called *complementary* or *antagonistic*. The complementary pairs of colours as above are roughly:—

Red and Green,  
Orange and Blue,  
Violet and Yellow,

a relationship that is indicated in the simple colour-circle, complementary colours being at the opposite ends of the three diameters, R.G., O.B., Y.V.

Colours and their relation to each other and to white may, however, be more accurately represented on an incomplete curve forming the two

<sup>1</sup> The simplest physical data concerning colour directly involve a physiological factor, inasmuch as the retina is the instrument of investigation. We shall therefore include theories of colour vision in this connection in order to avoid repetition. The subject is, however, more suitable for a second reading after the physiology of the retina has been studied.



sides of a triangle, than on a complete circle as above. To understand the construction of the colour-triangle, we may imagine the force or stimulation-energy of colour to be represented by weights. If each colour were an equal force, then each of the colours R. O. Y. G. B. V. would be represented by equal weights at equal intervals on the circumference of the circle, and the resultant of their fusion would be at W., their centre of gravity—*i.e.* the centre of the circle. But this is not the case; the component colours of white light are not of equal stimulation-energies, and may not therefore be represented by equal weights. These differences are taken into account in the construction of a colour-triangle, which gives the colours as if represented by weights proportional to their stimulation-energies, white as the resultant of their fusion being at the centre of gravity of the system. A properly constructed colour-triangle exhibits, moreover—(1) the relative strengths of the different colours, these strengths varying inversely as the dis-

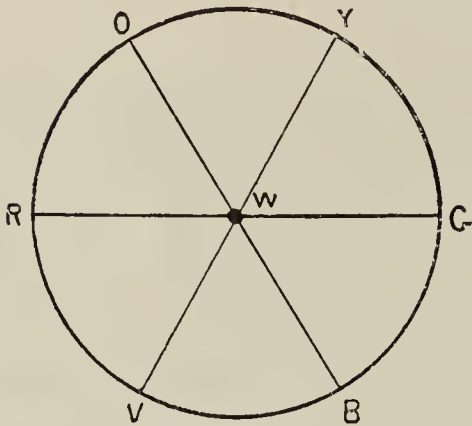


FIG. 214.—COLOUR-CIRCLE.

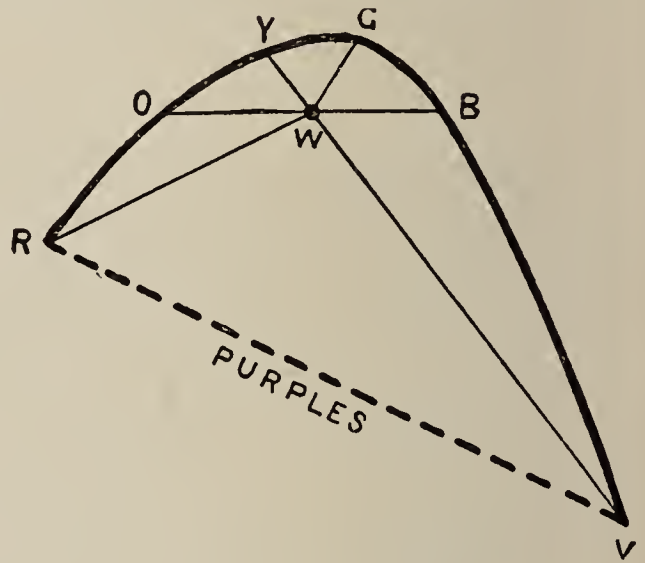


FIG. 215.—COLOUR-TRIANGLE.

tances from W.; (2) the position of complementary colours, these being situated at opposite ends of straight lines drawn through W. to the boundary of the triangle; (3) the position of mixture-colours, these being on the straight line joining the positions of the constituent colours—*e.g.* mixtures of the red and green being on the line R. G., of green and violet on the line G. V., of orange and blue on the line O. B.; (4) that the primary colours R., G., and V., being at the angles of the triangle, are not producible by mixtures of any of the other colours. The broken line forming the base of the triangle between R. and V. expresses the fact that the spectrum does not return upon itself, and cannot therefore be represented by a closed curve, but that a series of non-spectral purples exist in Nature, grading from red to violet, complementary to green and to the neighbouring colours at the apex of the triangle.

Differences of colour are of three kinds:—(1) Tone or wave-length. (2) Saturation or fulness. (3) Brightness or strength. It is easier to show by examples what is meant by these terms than formally to define them. We cannot, for instance, express the energy values of

two different colours in terms of a common unit, and we are very liable to confuse physical measures of the stimulus with physiological measures of the sensation.

The pairs of terms given above are those generally used as synonymous; but we may advantageously distinguish between the terms 'brightness' and 'strength,' for instance, as regards their application to light and to colour. We ought to apply 'brightness' to the *sensation*, 'strength' to the *stimulus*, just as we apply 'sweetness' to the sensation of a sugar solution, 'strength' to the stimulus as measured by weight of sugar. But we have first to distinguish between the three pairs of terms given above.

*Tone* or *colour* is the popular, or scientific, or artistic name by which different kinds of light are distinguished—red, green, violet, &c. The wave-length of a homogeneous colour defines its position in the spectrum. The wave-lengths of the spectral colours range from  $\frac{700}{1000000}$  at the red end to  $\frac{400}{1000000}$  at the violet end; at about the middle of the spectrum (green-blue) the wave-length is  $\frac{500}{1000000}$  or  $\frac{1}{2}\mu$ . Mixture tones may be recognised as such and their components identified by submitting them to the test of a prism—*e.g.* a homogeneous orange remains orange: a mixture-orange, indistinguishable from it by the naked eye, exhibits the red and yellow when viewed through a prism. *Saturation* or *fulness* is to be carefully distinguished from brightness or strength; it depends upon the amount of white light with which the colour may be mixed; a colour is full or saturated when it is free from white light, pale or unsaturated when it is mixed with much white light. *Brightness* or *strength* of colour depends upon the strength of the white light from which it is derived; a colour is dark or bright according as there is much or little of it.

It is desirable to recognise the terms in common use which it is correct or not correct to use in opposition to each other. It is not correct to use pale and dark as opposed terms, nor even light and dark as applied to colour; the accurate terms to use are 'pale' or 'light' *versus* 'full' as regards saturation, 'dark' *versus* 'bright' as regards strength. Pale or light red is a dilute red in much white, dark red is a weak red derived from a small quantity of white light. And the same for other colours, the common names of which may be grouped as follows:

Pale or 'dilute'	Normal	Dark or 'weak'
Rose or Flesh	Red	Maroon
Straw	Yellow	Brown
Light green	Green	Olive
Light blue	Blue	Dark blue

The distinction between saturation and brightness is very well illustrated by contrasting the effects of one of Maxwell's colour-discs—say red—in combination with a white disc, and in combination with a



black disc. In the first case the red is diluted with white, and a pale, or light, or less saturated red is the effect; in the second case the amount of red is reduced, and a dark red is the result. We may give one more example of the distinction. A red surface illuminated by white light may be said to reflect red rays and to absorb all others. A

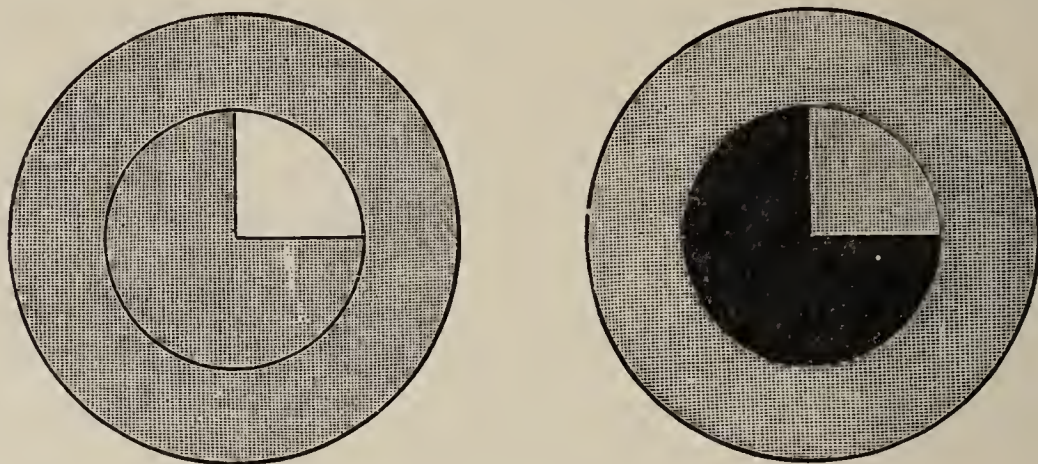


FIG. 216.—MAXWELL'S DISCS.

Arranged to illustrate difference of saturation and difference of brightness. In each case the outer circle is red; in the left-hand disc, the central part is three parts red to one part white, and when rapidly revolved appears *pale red*; in the right-hand disc, the central part is one part red to three parts black, and when rapidly revolved appears *dark red*.

highly saturated red surface illuminated by a strong white light is 'bright' red; if faintly illuminated it is 'dark' red. An unsaturated red surface illuminated by white light is 'light' or 'pale' red.

The distinction that we alluded to above as being desirable between brightness and strength, although not easy nor even possible to observe strictly, serves, however, to call attention to a distinction which is of the greatest importance, viz. that between sensory comparisons and stimulation measurements. We shall find in a later section (p. 554) that the ratio between these two variants is not arithmetical but geometrical, and that our measure of sensation is an indirect one by a varying unit. Differences of 'tone,' of 'brightness,' and of 'saturation,' are discriminated in sensation; these names are therefore applicable to the subjective aspect of colour-differences. Wave-length, and in less degree saturation, are quantities accessible to instrumental measurements; these terms are therefore applicable to the objective aspect of colour-differences. But we can draw no hard-and-fast line in the terminology; the strength of a stimulus is measurable, but we often speak of the strength of a sensation which can be expressed only in ratios, and not in fixed units. Saturation is indiscriminately applied to the objective and to the subjective aspect of colour-quantity. Physically, all the spectral colours are saturated; physiologically, some spectral colours are more saturated than others. A physically saturated colour is a maximum value which may be represented by a fixed number; the sensation to which it gives rise is its physiological maximum effect, but the latter is not a constant, and cannot be represented by a fixed number; the fixed maximum stimulus of spectral red will give

rise to a smaller physiological maximum by an eye which is fatigued by red, and to a larger physiological maximum by an eye which is fatigued by the complementary green-blue. We shall not pursue these considerations further, having said enough to remind the reader that the subjective factor is variable, and that it plays a large part in the estimation of colour-quantities—necessarily so, since the instrument by which we can measure and compare differences of that form of radiant energy called light, or colour, is the retino-cerebral apparatus by which we feel them.

*Absorption of light.*—Opaque bodies arrest the passage of all light; transparent bodies, like glass or water, allow light to pass unchanged provided they are themselves colourless. But coloured transparent bodies stop the light of certain parts of the spectrum, while they allow that of other parts to pass. Thus red, yellow, blue, or otherwise coloured glass allow only red, yellow, blue, or other coloured light to pass, the remainder of the spectrum being in each case stopped by absorption. That this is really the case—*i.e.* that the light which passes through a red glass is white light *minus* orange, yellow, green, blue, and violet, and not white light *plus* red light—may easily be verified by looking at the spectrum of white light passed through a red glass; only the red end of the spectrum remains visible; the light on the violet side of the D line is stopped, or, as it were, ‘filtered off,’ by the red glass.

The colours of bodies viewed by reflected light are likewise due to absorption. Red velvet illuminated by white light absorbs all except red light, which it reflects to the eye.

It is not always possible to tell by inspection of a coloured body what will be the precise portion of the spectrum that will be absorbed by it; in the familiar instance of blood, for example, the absorption bands revealed by the spectroscope are such as could not have been foreseen from the colour of diluted blood, and are so peculiar as to afford a delicate means of identification. The same example may be utilised to show that the absorption which takes place with reflected light is identical with that which takes place with transmitted light; if a little blood be smeared over a piece of white paper, and looked at with a spectroscope in a strong light, the absorption bands are seen just as if a layer of blood were looked *through* in the ordinary way. The last instance is, indeed, such as to indicate to us that the absorption of reflected light is essentially the same as that of transmitted light, inasmuch as the coloured reflected light is actually transmitted *through* a superficial film of the body that is looked at.

Coloured papers or other surfaces reflect a more or less extensive portion of the spectrum—*e.g.* a yellow paper reflects some red and green as well as yellow, a blue paper reflects green and violet as well as blue, a purple paper reflects red and blue, as can be verified by means of a spectroscope. By the same means we can distinguish whether a



prismatic colour is simple or compound; a blue spot, for instance, might be caused by a spectral blue, or by the superposition of spectral green and violet; the first blue would, *cæteris paribus*, be more saturated than the second, and on viewing the latter through a spectro-scope its compound nature would become apparent.

*Retinal fusion of colours.*—1. A vertical plate of glass placed between two horizontal coloured surfaces—*e.g.* blue and yellow—reflects one colour, while the second colour is seen through the glass. The

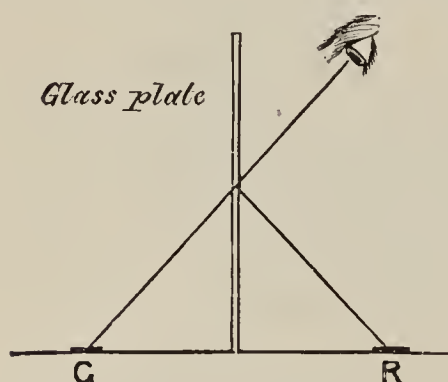


FIG. 217.

two colours are thus simultaneously viewed together, and their resultant in sensation is a single effect approximating to white.

2. A rapidly-rotating disc upon which sectors of any two or more colours are fixed, gives in sensation a resultant which is a single compound colour. Each colour is viewed in rapid succession, and, owing to the persistence of retinal impressions, the two or more constituent colour-impressions cover each other and blend, giving a single compound colour. This is the method alluded to above as that by Maxwell's discs.

3. Two separate prismatic spectra are superposed so that various colours coincide. With suitable apparatus this method is by far the best; the separate colours are the purest possible, and the resultants from their mixture are correspondingly pure. The first two methods are only suitable for demonstration, for colours painted upon a surface are necessarily impure.

*Physiological theories.*—Most of the facts relating to colour-vision, its defects, and various experimental modifications, may be accounted for in accordance with a theory known as the *Young-Helmholtz theory*. It is as follows: The cones of the retina, or the nerve-fibres connected with them, are of three different kinds as regards their excitability by various colours of light; one kind of cones is most excitable by red rays, less excitable by green and violet rays; a second kind is most excitable by green rays, less excitable by red and violet rays; a third kind is most excitable by violet rays, less excitable by red and green rays. This supposition is represented by the three curves R, G, V in fig. 218.

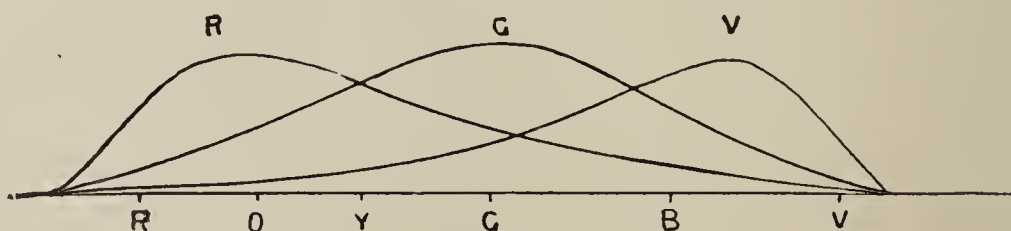


FIG. 218.

This theory is to be regarded more as a convenient standard to which various facts may be referred, rather than as itself a material demonstrated fact. It should not be employed as an explanation of facts, but only as a convenient supposition upon which certain facts

may provisionally be placed together.<sup>1</sup> That the theory is not an actual expression of fact is clearly indicated by the following experiment:—A spot of white light forming on the retina an image less than  $2\mu$  in diameter does not assume any colour if cast upon various points, effects of chromatic dispersion being excluded; a minute spot of homogeneous colour does not alter under similar conditions; on the contrary, colour differences are more and more difficult to distinguish the smaller the area from which they proceed, and if exceedingly minute are no longer distinguished at all, but seen simply as colourless light.

The alternative theory is that of Hering. The visual apparatus is supposed to be capable of six fundamental distinct sensations—white and black, red and green, yellow and blue. White, red, and yellow sensations are katabolic—*i.e.* are concomitants of disintegration; black, green, and blue are anabolic—*i.e.* concomitants of integration. The visual substance is further supposed to be of three kinds: changes of the first give white or black; of the second, red or green; of the third, yellow or blue. The white-black substance is supposed to be influenced by the whole range of the spectrum; the red-green to be used by red rays, restored by green rays; the yellow-blue to be used by yellow rays, restored by blue rays.

The essential idea common to Hering's as well as to Helmholtz's theory is, that the sensation of light is the resultant of sensificatory antecedents of *three* different kinds in varying proportions—*i.e.* that the sensation is the resultant of a three-dimensional stimulus. But Hering's three dimensions differ from Helmholtz's.

Wundt, admitting the physical fact that any kind of light sensation can be produced from three fundamental colour-stimuli, rejects

<sup>1</sup> The purely imaginary nature of this hypothesis is insisted upon by Helmholtz himself, who, in the second edition of his *Physiological Optics* (1889, p. 349), supplements it by the assumption of three kinds of photo-chemically changeable substances, and of three kinds of cells in the cerebral cortex. Hering expresses himself as follows: 'The psycho-physical vision-stuff may accordingly be regarded as a mixture of three qualitatively different substances. Nevertheless, I signified long ago (*Zur Lehre vom Lichtsinne*) that I gave this form to the theory only for the sake of greater distinctness, and that I consider it to be fundamentally more correct to say that the psycho-physical vision-stuff is capable of qualitatively different kinds of Dissimilation and Assimilation, and that each of these innumerable different kinds of D. and A., assuming three principal kinds of D. and A., can be conceived as broken up into three components, each of which corresponds to one of these three principal kinds of D. and A. It is, no doubt, more intelligible to most people to say that each visual sensation may be conceived as composed of given primary or elementary sensations in given proportions, than to say that each visual sensation takes, in the pre-arranged three-dimensional variability of visual sensation, a given place in relation to three given sensation series; and it is more intelligible to regard the vision-stuff as a mixture of three separate substances, each of which can be altered in two qualitatively opposite senses (D. > A. and A. > D.), than to say that the variability of state of vision-stuff is three-dimensional.'



the three-component theory, and holds that retinal excitation by light gives rise to two different kinds of excitatory process, viz. (*a*) an achromatic process, (*b*) a chromatic process, varying in degree with intensity and with wave-length, the former being relatively more efficacious than the latter with very weak and with very strong stimulation. He urges that a sensation of light and dark must be antecedent in development to a sensation of different colours of light, and that we cannot regard the achromatic sensation as a normal product of chromatic factors.

*Colour-blindness.*—A considerable percentage of people (5 per cent.) fail to distinguish colours from each other which to the majority of people are perfectly different. When this inability is very marked its subject is said to be colour-blind. Colour-blindness varies in degree and character, but by far the commonest form is that in which the discrimination of red from green is imperfect. This defect has very obvious practical importance as regards railway and steamship officials, who should be able to distinguish without hesitation between green and red lights. The assured fact that a given person confuses red and green—*i.e.* that red and green rays excite the same kind of sensation—is attributed to the by no means assured fact that cones most excitable by red rays are not present in his retinae (Helmholtz), or that the red-green visual substance is wanting (Hering).

*The colour-perimeter.*—If, while the eye is fixed upon a point directly in front of it, the extent of the field of indirect vision be tested, by moving a small disc so as to come in and out of the field from several sides, it will be found that the extent of the field of indirect vision is greatest for a white disc, smallest for a red disc, and that the field is greater for green than for red, and greater for blue than for green. (See also p. 431.)

## VISION

The organ of vision is composed of (1) a *receptive* part—the eye; (2) a *transmissive* part—the optic nerve; and (3) a *perceptive* part—the cortex of the brain over the occipital lobes.

The eye is essentially a *camera obscura*, having anteriorly a *biconvex lens* by which real reversed <sup>1</sup> images of external objects are focussed upon a sensitive surface, *the retina*. Muscles attached to the eyeball turn the eye in various directions towards various points of the field of vision. The exact focussing of images formed by objects near to and far from the eye is effected by alterations of the focussing power of the lens, and not, as in the photographic camera, by alterations of the distance between lens and sensitive

<sup>1</sup> A peculiar exception to this rule occurs in the compound eyes of insects, where the formation of a real *erect* image has been demonstrated by Exner.

surface. The focussing power of the lens is altered by the action of the ciliary muscle—this change constituting the act of accommodation. A circular diaphragm—the iris—cuts off light from all but the central part of the lens, and contracts or dilates by muscular action under a variety of conditions.

The eye is almost spherical; the greater part of its wall is composed of three coats—the sclerotic, choroid, and retina. The posterior  $\frac{5}{6}$ ths of the wall are opaque, the anterior  $\frac{1}{6}$ th of the wall is transparent and is named the cornea; it is continuous with the sclerotic. The iris is continuous with the choroid; both usually

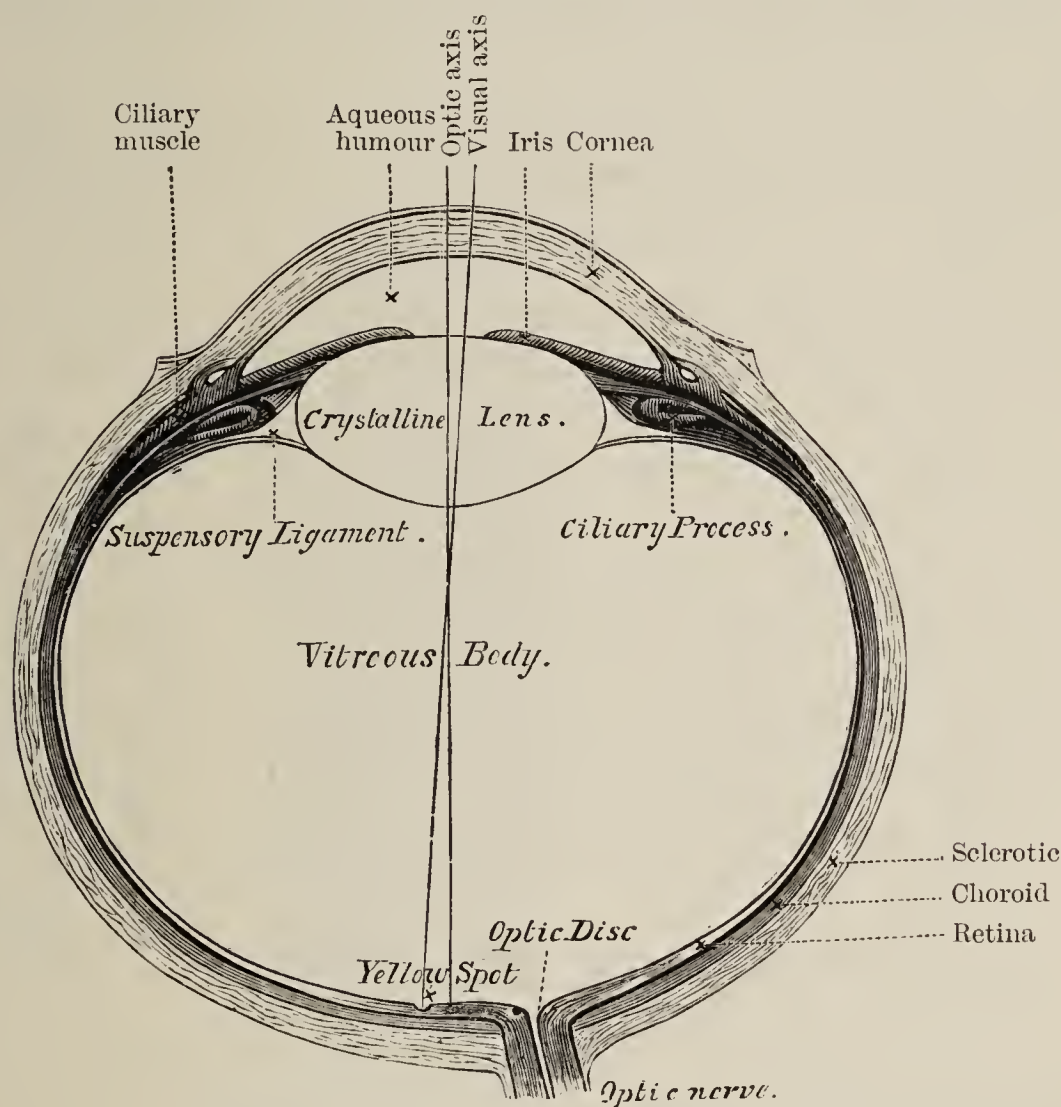


FIG. 219.—HORIZONTAL SECTION OF THE LEFT EYEBALL.

are deeply pigmented, and effectually convert the interior of the eye into a dark chamber; the pupil, or centre aperture of the iris, is made smaller or larger by the contraction of circular muscular fibres in the iris (constrictors), or of radiating muscular fibres (dilators). The circular or constrictor fibres are excited to contraction through the channel of the third or motor oculi nerve; the radiating or dilator fibres are excited to contraction through the channel of the cervical sympathetic. The retina is the expansion of the optic nerve as the internal of the three coats; it occupies the posterior  $\frac{2}{3}$ ths of the internal surface, and is



composed of several layers. The most external of these layers—the layer of rods and cones—is excitable by light, and is the true end-organ of the optic nerve; it is, like all other sensifactory surfaces, composed of elements which are modified epithelial cells, originally derived from the external epithelium of the embryo (the epiblast). The contents of the eyeball are—(1) the aqueous humour in the anterior chamber, (2) the crystalline lens held in place by its capsule and suspensory ligament, to which is attached the ciliary muscle, (3) the vitreous body. These three substances are transparent, and with the cornea constitute the refractive media of the eye, which conjointly act as a converging lens.

The retina is a comparatively thick stratified membrane, comprising from vitreous to choroid surface the series of layers shown below.

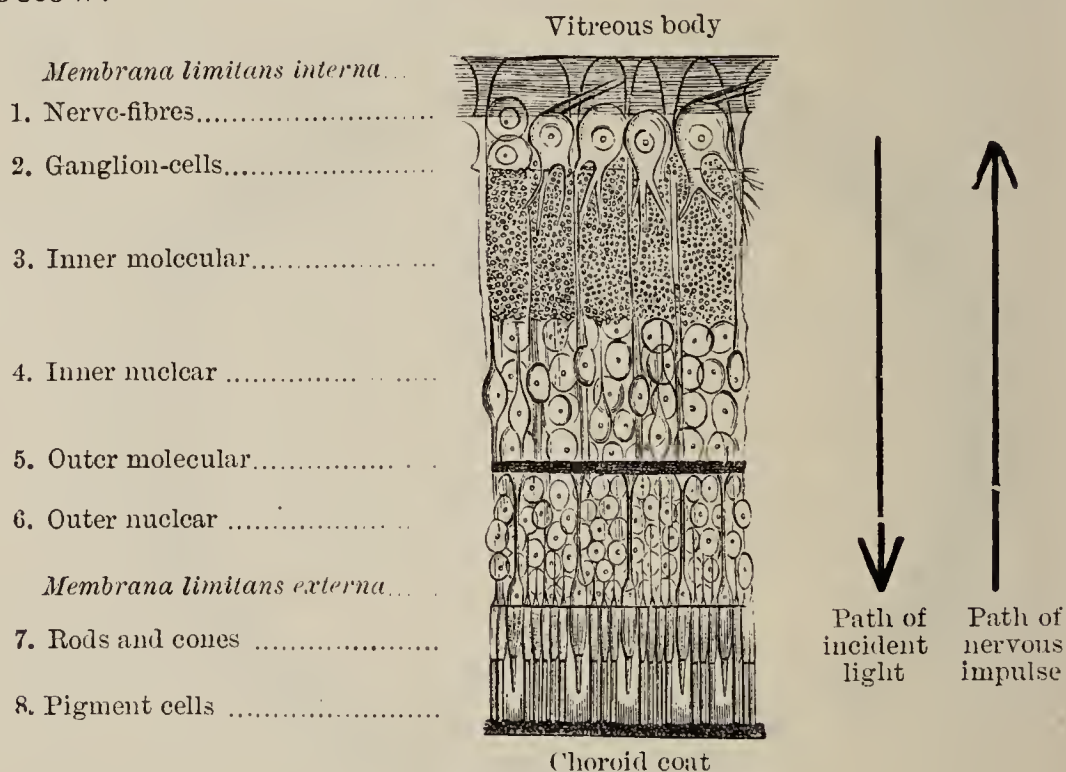


FIG. 220.—THE RETINA.

The elements composing these several layers are held together by 'sustentacular' connective tissue, which supports the nervous elements proper and forms membranous sheets—the external and internal limiting membranes—between which are included the inner first six layers of the retina.

The essential terminal apparatus is constituted by the rods and cones of the seventh layer, which is contiguous with the choroid, and not, as might have been imagined, the superficial layer first exposed to the impact of light. A ray of light must traverse the layers in the direction from 1 to 7 before it gives rise to a nervous impulse at the seventh layer; this impulse travels through the layers in the reverse direction, from 7 to 1, and is led off by the fibres (non-medullated) of this layer, which

converge to form the optic nerve, where they become medullated. The retinal blood-supply is derived from the central artery, which branches out from the optic disc and forms a superficial ramification, the capillaries from which are confined to the first four layers of the retina; the remaining layers are therefore extravascular; and it is to be noticed that the vessels are thus situated on the path of light in front of the layer of rods and cones, so that it is possible for shadows of them to be cast upon this layer.

The structure of the retina differs from that indicated above at two exceptional points, viz. at the optic disc and at the yellow spot. The optic disc is the point at which the optic nerve enters the eyeball, and whence its fibres spread out in all directions over the inner surface of the eyeball. This spot is destitute of true retina and is not excitable by light; its situation and dimensions can be experimentally determined by Mariotte's experiment. At the yellow spot, or, more precisely, at the fovea centralis, or central depression in the middle of the yellow spot, the retina is very thin, consisting of little more than a single layer of attenuated cones. It is the region of most distinct vision—*i.e.* the spot upon which objects are imaged when they are looked at; but *not* the area of the greatest sensibility—on the contrary. These anatomical points, taken in conjunction with the experiments of Mariotte (p. 423) and of Purkinje (p. 452), constitute proof positive that the rod-and-cone layer of the retina is the layer excitable by light and excitatory of visual sensation.

It is to be remembered that the eye as a converging lens differs from an ordinary biconvex lens in this respect, that the rays are twice refracted at the surfaces of the latter—once at entrance from air, once at exit into air; whereas in the eye the chief refraction occurs at the entrance-surface from air, namely, at the cornea, subsequent refraction by the other media being of comparatively small amount. Of these subsequent media, the crystalline lens is certainly the most important, as by it is obtained the additional refractive power required in accommodation. But that the lens is not absolutely essential to vision is clearly shown by the condition of sight after it has been extracted for cataract; accommodation is then lost, and the eye is hypermetropic by the loss of about 10 diopters lens power, but the patient obtains a useful eye. The power of such an eye requires to be supplemented by a 10-diopter glass for distant vision, and



for vision at a distance of 20 centimeters by a 15-diopter glass.

The various refractive media (of which the list on p. 424 gives the normal values) may, as regards their total converging effect, be considered equivalent to that of a single substance with a re-

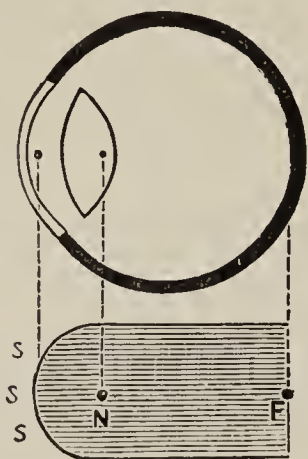


FIG. 221.—THE REDUCED OR SCHEMATIC EYE.

Radius of curvature of refracting surface, SSS	}	=	5 mm.
Index of refraction			
Position of nodal point, N	}	=	{ 5mm. behind SSS.
Position of principal focus, F			
	}	=	{ 15 behind N or 20 mm. behind SSS.

Such an eye would be approximately represented by a piece of glass shaped like the lower figure.

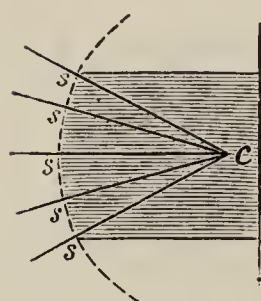


FIG. 222.

$s s s s s$  is the spherical surface of a mass of glass;  $c$  is the centre of curvature of that surface. The radii,  $c s$ , are perpendicular to the surface, and rays of light entering the glass on these lines will not be deflected, but will meet at the point  $c$ . Lines of this kind (viz. perpendicular to the surface of separation) are termed '*lines of direction*,' and  $c$ , their point of intersection, is termed the '*nodal point*.' In other words, the nodal point may be defined as the point of intersection of rays perpendicular to the refracting surface. In the case of the eye there are several such nodal points behind its several refractive surfaces. But in the schematic eye described above we have considered these surfaces and points as represented by one surface and one nodal point.

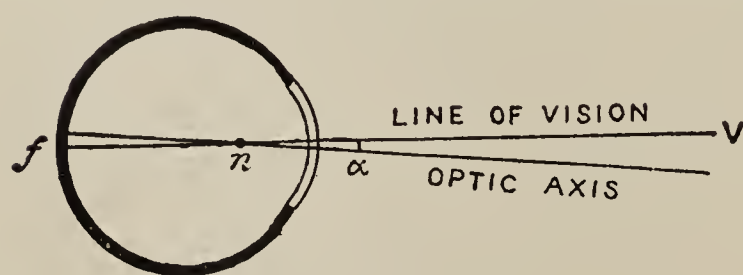


FIG. 223.

retina is between the fovea and the entrance of the optic nerve.

fractive index of 1.35, and a single spherical surface of 5.1248 mm. radius. In such a medium the distance between the nodal point,  $N$ , and the principal focus,  $F$ , is about 15 mm. Its collecting power is equal to about 50 diopters—*i.e.* it is a convex lens with a focal distance of 2 cm. This is a schematic or *reduced eye*. Taking the retina as the point of departure, the nodal point,  $N$ , of such a reduced eye, as compared with the normal eye, will be just in front of the posterior surface of the lens, and the one refracting

The *optic axis* is a straight line through the centres of curvature of the cornea and lens, prolonged to the posterior wall of the eye; the point at which the optic axis meets the

The *line of vision* may be defined as a straight line,  $f n V$ , between the fovea and any point to which the gaze is directed. It does not coincide with the optic axis, but forms with it an angle of about five degrees at the nodal point (fig. 223).

*Visual angle.*—The angle  $B n A$  included between two straight lines from the borders of an object  $B A$  to the nodal point  $n$  is called the visual angle; it is equal to the opposite angle  $b n a$ .

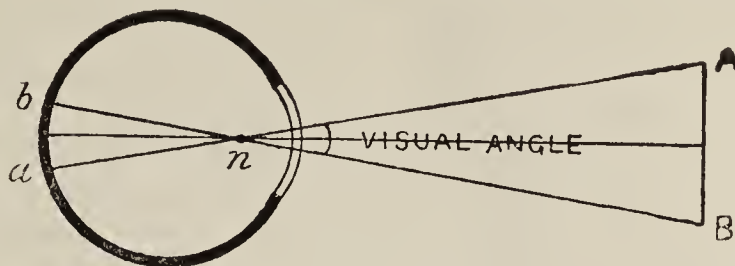


FIG. 224.

The value of the distance  $N F$  (fig. 221: 15 mm. in the reduced eye) supplies a necessary datum in calculating the size of the retinal image of an object—given the size of the object and its distance from the eye.

For example, what will be the height on the retina of an observer, of a man six feet high at a distance of one mile?

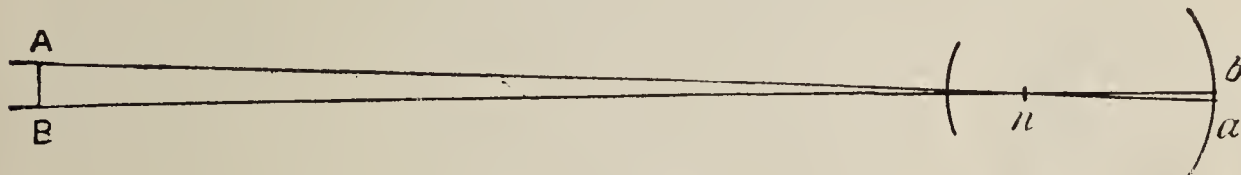


FIG. 225

Let  $A B$  represent the man,  $a b$  the image of the man on the eye of an observer,  $n$  being the nodal point of that eye.

From the two similar triangles  $A n B$ ,  $a n b$ , we have  $\frac{a b}{a n} = \frac{A B}{A n}$ , or, substituting the numbers given above,  $\frac{a b}{15 \text{ mm.}} = \frac{6 \text{ feet}}{1 \text{ mile}}$ ; that is,  $a b = 0.017 \text{ mm.}$ , *i.e.*  $17 \mu$ .

This is about twice the diameter of a blood-corpuscle. We know that a man will be visible at a much greater distance than one mile—*i.e.* when forming on the eye a retinal image much smaller than  $17 \mu$ —a consideration which brings home to the mind that it is, under certain conditions, possible to see blood-corpuscles in one's own retina.

Another good example of the application of the reduced eye, especially as regards the magnitude  $N F = 15 \text{ mm.}$ , is afforded by the following exercise:—Determine on your own eye the distance between the yellow spot and the entrance of the optic nerve—*i.e.* the blind spot (Mariotte's experiment). The problem can easily be solved as follows:—Pin a large sheet of white paper against the wall, mark on it a small black cross,  $C$ , on a level with the eyes; steadily look with one eye at the cross, having fixed the position of the head by a ruler of convenient length (say 50 cm.) held between the teeth and resting against the wall; while the gaze is thus fixed move a white quill pen with inked point towards the temporal side of the field of vision, and mark on the paper the exact spot at which the black point is lost sight of.



Under the above conditions the retinal image of the cross is on the centre of the yellow spot, and the second mark shows where the image of the black mark has passed over the temporal border of the blind spot.

In the experiment just made the distance between the two points A and C was 10 cm., that from *n* to C was 50

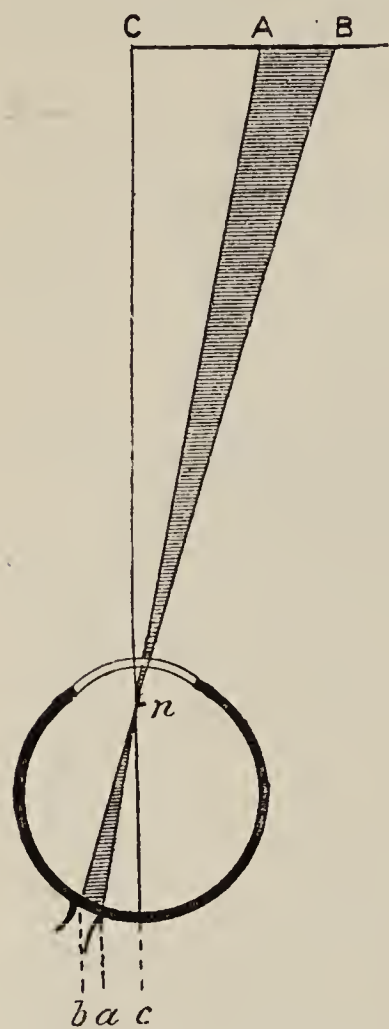


FIG. 226.—HORIZONTAL SECTION OF THE RIGHT EYE, TO ILLUSTRATE MARIOTTE'S EXPERIMENT.

cm.; therefore,  $\frac{10}{50} = \frac{ac}{15}$ ; that is, *ac*, the distance between the temporal border of the optic disc and the centre of the yellow spot = 3 mm.

Similarly, by moving the black-tipped pen in various directions we may determine the limits of the blind field, and from its magnitude, *AB*, deduce the retinal magnitude of the blind spot, *ab*. In the above example the horizontal magnitude, *AB*, was 6 cm.—*i.e.* the retinal magnitude was 1·8 mm.; measured vertically, the diameter of the blind field was 8 cm.—*i.e.* its retinal magnitude was 2·4 mm.

To determine the smallest perceptible image on the retina.—Gum two slips of white paper 1 mm. wide parallel with each other on a black card, leaving an interval 1 mm. broad between the two slips. Determine the distance from the eye at which the two white slips are just visible, or just invisible as a *double* line. In the observation just taken the distance was about 4 meters.

As before,  $\frac{ab}{an} = \frac{AB}{An}$  (fig. 225), or, substituting the appropriate numbers,  $\frac{ab}{15 \text{ mm.}} = \frac{1 \text{ mm.}}{4,000 \text{ mm.}}$  *i.e.* *ab* = ·00375

mm., or 3·75  $\mu$ . The result agrees with the classical results of Helmholtz and others, according to which the smallest angular distance at which points can be separately distinguished is 50 sec., with which the size of a retinal image is 3·65  $\mu$ . It is to be observed that this magnitude closely coincides with the diameter of cones at the fovea, which is about 3  $\mu$ , the distance between the centres of adjacent cones being about 4  $\mu$ .

AVERAGE MEASUREMENTS OF THE NORMAL EYE

					mm.
Antero-posterior diameter of eyeball	.	.	.	.	24
Radius of curvature of cornea	.	.	.	.	8
"	"	ant. surface of lens (non-accom.)	.	.	10
"	"	"	"	(accom.)	6
"	"	posterior surface of lens	.	.	6
Refractive index of aqueous humour	.	.	.	.	1·33
"	"	vitreous humour	.	.	1·33

	mm.
Mean refractive index of lens . . . . .	1.45
Diameter of cornea . . . . .	12
Distance between cornea and lens . . . . .	4
Mean diameter of pupil . . . . .	4
Thickness of cornea . . . . .	1
„ lens (non-accommodated) . . . . .	4
„ „ (accommodated) . . . . .	4.4
Thickness of retina at fundus . . . . .	0.2 to 0.4
Distance between retinal vessels and rod-and-cone layer . . . . .	0.2 to 0.3
Diameter of optic disc . . . . .	1.5
„ yellow spot . . . . .	1.25
„ fovea centralis . . . . .	0.25
„ cone in fovea . . . . .	0.004 to 0.005

The total number of cones is estimated to be 7 millions, of rods 130 millions, of fibres in the optic nerve,  $\frac{1}{2}$  to 1 million.

**Accommodation.**—The formation of a distinct image upon the retina is necessary to distinct vision. With any given lens, distinct images of different objects, near or far, may be obtained by altering the distance between the lens and the surface upon which the image is to be formed; if this distance is unalterable, a more or less convergent lens must be used for near and for distant objects respectively. In the case of the eye the distance between the lens and the retina is invariable, but the lens may be rendered more or less convex by more or less muscular action. The lens is lodged between two layers of the suspensory ligament, which is a prolongation of the choroid, and is kept compressed by the tension of that membrane; a circular muscle, the *ciliary muscle*, having its fixed point at the junction of the cornea and sclerotic, and its insertion in the choroid, can pull upon the choroid so as to relax the tension of the suspensory ligament, when the lens by virtue of its elasticity becomes more convex. A more convex lens is a more convergent lens, and has a shorter focus—*i.e.* divergent rays from a near object which in a resting eye with compressed lens would be focussed behind the retina are now brought to a focus on the retina itself, and the object is seen distinctly. The change thus effected by the agency of the ciliary muscle constitutes the act of *accommodation*, which is a



FIG. 227.

To illustrate accommodation: non-accommodated compressed lens represented clear; additional curvature by the bulging of its anterior surface in extreme accommodation represented by the shaded crescent.



voluntary act (although the ciliary muscle is composed of smooth fibre). With relaxed accommodation the eye is adjusted to distinct vision of distant objects, the rays from these to the eye are practically parallel, the suspensory ligament is tense, the lens is compressed, the ciliary muscle is at rest. With accommodation the eye is adjusted to distinct vision of near objects,

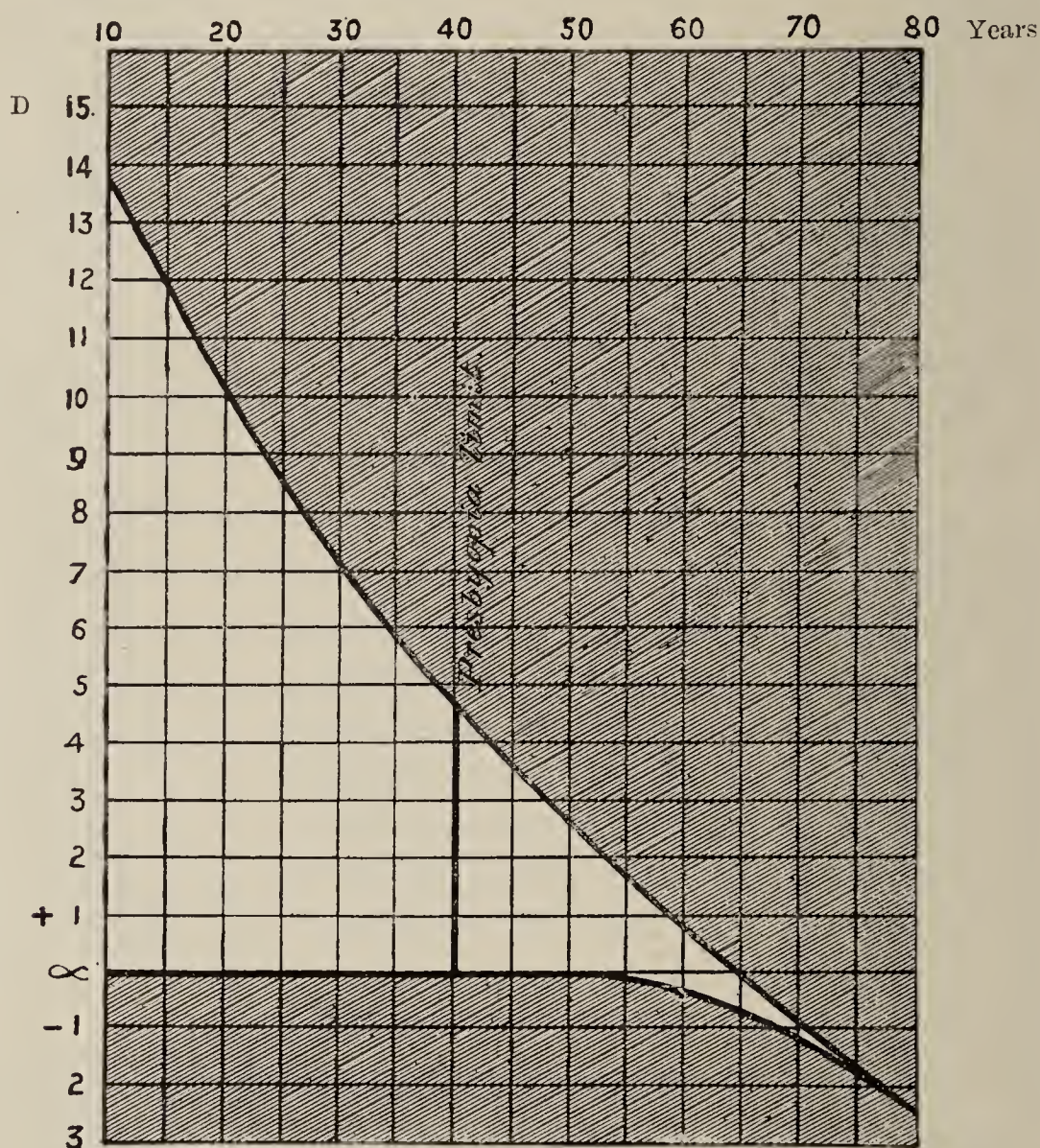


FIG. 228.—POWER OF ACCOMMODATION AT DIFFERENT AGES. (Donders.)

The ordinates indicate power in diopters. Age is indicated by the numbers along the abscissa—*e.g.* at 20 years the range is between infinity for the resting eye and 10 diopters for the accommodated eye; at 30 years the range is from infinity to 7 diopters; soon after 50 years the resting eye is under-focussed for infinity; soon after 65 years, even a fully accommodated eye is under-focussed for infinity. The focussing power must be supplemented by convex glasses, increasing in strength with increasing age.

the rays from these to the eye are divergent, the suspensory ligament is less tense, the lens is less compressed, the ciliary muscle is in action. With the normal eye distant objects are seen without accommodation, the converging power of the lens and the length of the eye are such that distinct images are formed on the retina of objects from which the rays to the eye



are parallel. Accommodation is only required for near objects, and in the normal eye the power of accommodation is such as to procure distinct images of objects up to about 5 inches from the eye; within this distance distinct images can no longer be obtained. The most distant point of which distinct vision is possible is called the 'far point'; the nearest point of which distinct vision is possible is called the 'near point'; in the normal eye of a person 30 years old the far point is at an infinite distance, the near point is at about 5 inches from the eye.

FIG. 229. — RECESSION OF PUNCTUM PROXIMUM WITH INCREASING AGE.

The ordinates in millimeters indicate the distance of the 'near point' in centimeters. When the 'near point' is further than 22 cm. presbyopia is said to begin. Normally, this occurs at the age of 40. In more or less myopic eyes the 'near point' is at about 10 cm., the 'far point' at 30 to 50 cm. The 'far point' of the normal eye is at infinity.

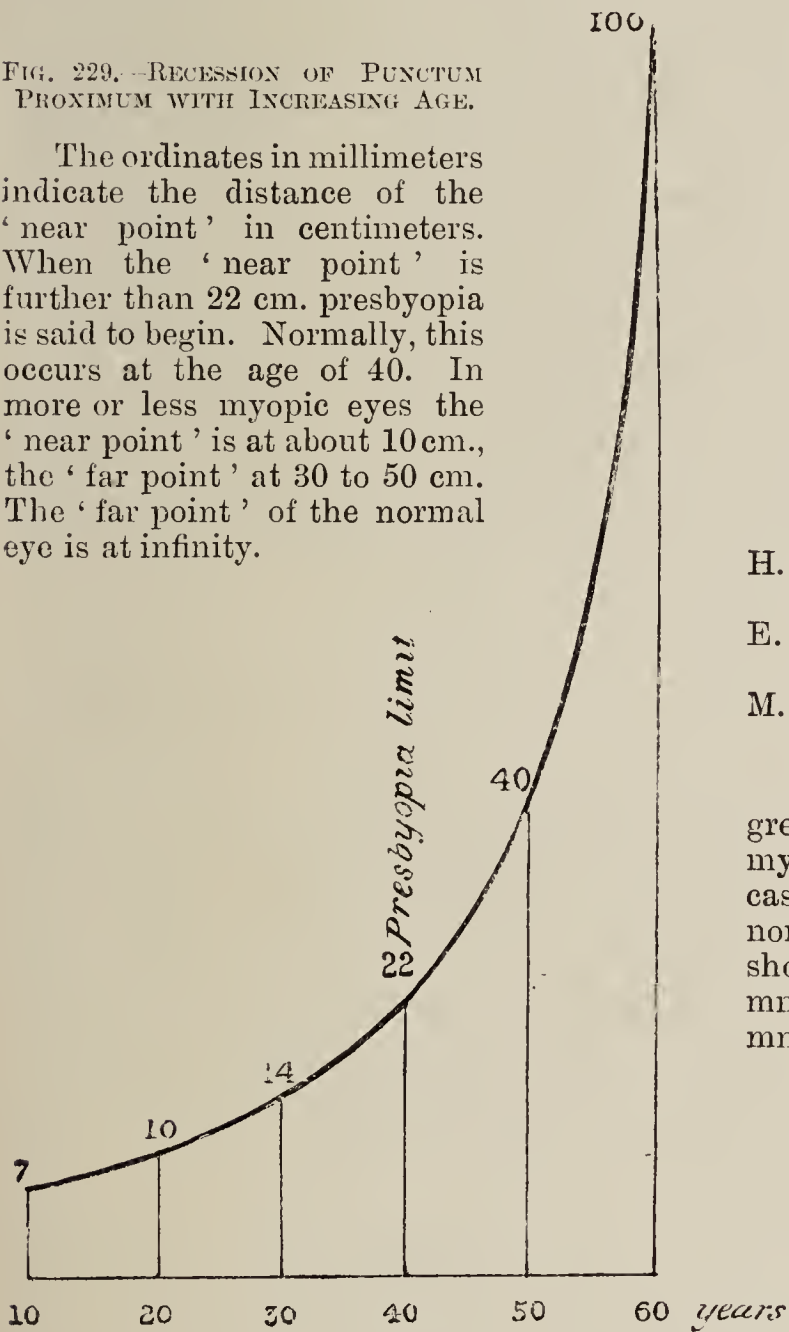


FIG. 230.

- H. *Hypermetropia*. — Short eye. Long sight.
- E. *Emmetropia*. — Normal eye. Normal sight.
- M. *Myopia*. — Long eye. Short sight.

The figure represents high degrees of hypermetropia and of myopia, *i.e.* an alteration in each case of about 12 diopters. The normal eye being 22 mm. long, the short eye will then be about 18 mm., and the long eye about 26 mm. in length.

*Myopia*.—The vision of distant objects is indistinct, and objects can be distinctly seen nearer than usual—the subject is said to be 'short sighted.' The defect is due to the fact that the converging power is too great in relation to the length of the eye, or, more correctly, that the eye is too long in relation to its converging power; parallel rays are focussed in front of the retina, and even very divergent rays can be focussed on the



retina. A scattering or biconcave lens is the corrective of this too great converging effect.

*Hypermetropia and presbyopia.*—Near objects cannot be distinctly seen, distant objects may be clearly seen. The subject is said to be ‘long sighted.’ The eye is not long enough in relation to its converging power; divergent rays cannot be focussed on the retina, but have their focus behind it; parallel rays may be brought to a focus on the retina, unless the affection be excessive. The hypermetropic eye is shorter than the normal eye, an effort of accommodation is requisite even to distant distinct vision, and if the hypermetropia be excessive even this is insufficient to effect focussing upon the retina. The *presbyopic* eye has a rigid lens or a weak ciliary muscle, so that the lens cannot become convex when accommodation is attempted. A gathering or biconvex lens is the corrective of these deficient converging effects, both in hypermetropia and in presbyopia.

Taking the normal non-accommodated eye to be represented by the schematic eye 2 cm. long, focussing on its posterior surface parallel rays from infinity, *i.e.* equal to a lens of 50 D, it is evident that an additional power will bring divergent rays to a focus on that surface, as follows :—

	+ 15	14	13	12	11	10	9	8	7	6	5	4	3	2	1	Diopters
will focus rays } from a point }	6·6	7·1	7·7	8·3	9	10	11	12·5	14·3	16·5	20	25	33	50	100	cm. distant.

A review of this table or of fig. 228 suggests points of great practical importance as regards the eyesight during adolescence. A child has great power of accommodation, and may readily form the habit of reading or writing too close to his book; this same power enables him to compensate an unsuspected hypermetropia that may be present by strong or, it may be, spasmodic accommodation, a habit that, if unchecked, causes headache and, in later life, defective accommodation.

*Sanson's images.*—That it is the lens and lens only that is altered in the act of accommodation is proved by the observation of ‘Sanson’s images.’ A candle held in an appropriate position close to an eye gives three reflections, which can be seen by an observer placing himself at an appropriate angle or more conveniently with the aid of the instrument known as the ‘*phakoscope*.’ These are reflections from the mirror surfaces :—1, of the cornea; 2, of the anterior surface of the lens; 3, of the posterior surface of the lens. The first image is very bright and unmistakable; it is a reflection from the convex surface of the

cornea, and is therefore an erect virtual image. The second image is larger and much less distinct; it is a reflection from the convex anterior surface of the lens; this surface is less convex than that of the cornea, and the image is erect, and larger than the corneal image. The third image is intermediate in distinctness between the first two, and very small; it is a reflection from the concave posterior surface of the lens, and is therefore a real reversed image. If, while the observer has in view these three images (or, better still,

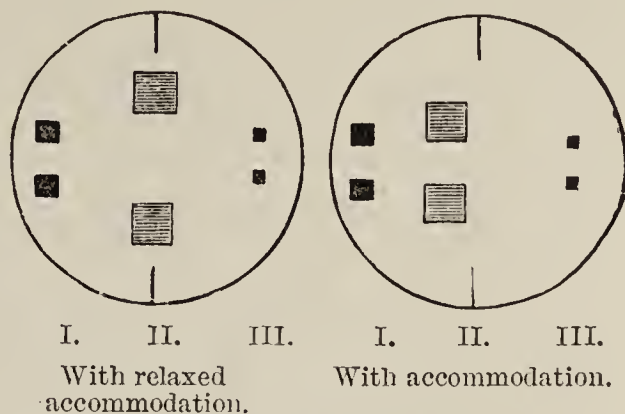


FIG. 231.—SANSON'S IMAGES.

- I. From anterior surface of cornea.
- II. From anterior surface of lens.
- III. From posterior surface of lens.

three pairs of images from a double light), the subject of observation alternately relaxes and accommodates (*i.e.* looks alternately far and near in one line of vision), it may be seen that images I. and III. remain immobile and unchanged, but that image II. advances towards image I. and becomes smaller with the act of accommodation; it recedes again from I. and recovers its size with relaxed accommodation. The stability of images I. and III. in the act of accommodation proves that the cornea and posterior surface of the lens do not move nor alter their curvature; the diminution in size and advance of image II. prove that the anterior surface of the lens becomes more convex and approaches the cornea.<sup>1</sup> This testimony is proof that the act of accommodation is accompanied by increasing convexity of the anterior surface of the lens, and of the anterior surface of the lens only.

The fact may be observed, but not so accurately, by directing a person to look near and far along one line of vision, while his eye is looked at from the side so that the black border of the pupil is just visible. The iris may be seen to be pushed forwards with the act of accommodation. This is, of course, only a rough and partial observation of the fact, which is proved with great nicety by means of the three images.

In the course of these observations it will hardly have escaped notice that the pupil is larger when the eye is adjusted for distant vision, smaller when the eye is accommodated to near

<sup>1</sup> Image III. does alter a little, but not enough to sensibly invalidate the statement made above. A fourth image, produced by the posterior surface of the cornea, is visible under favourable conditions.



vision—*i.e.* contraction of the pupil accompanies the act of accommodation.

The *field of vision* is the entire surface from which rays of light reach the retina; in other words, it is the total visible area in front of the open eye. The term is somewhat loosely applied with reference to the moving or to the stationary eye; strictly speaking, it applies to the latter only, and alters with every movement of the eye, its centre being always the particular spot from which rays are focussed upon the fovea centralis. This spot or point of fixation is the centre of a small area of distinct vision, which, again, is surrounded by a much larger area of indistinct vision. The area of distinct vision is included by an angle of about  $1^\circ$  only (the diameter of the fovea being only  $\frac{1}{4}$  mm., the diameter of the distinct area in the field of vision at a distance of 30 centimeters will be only  $\frac{1}{2}$  centimeter, at a distance of 30 meters only  $\frac{1}{2}$  meter), while that of indistinct vision is for each eye about  $160^\circ$  horizontally and  $120^\circ$  vertically; with both eyes open the horizontal range is over  $180^\circ$ .

To look at an object means to turn the eye in such a position and to accommodate its focussing power to such an extent that a well-defined image of the object falls on the fovea; we then have distinct or 'direct' vision of the object, and at the same time indistinct or 'indirect' vision of neighbouring objects, which form more or less well-defined images on the retina farther and farther from the yellow spot. The retinal image is thus 'like a picture minutely and elaborately finished in the centre, but only roughly sketched-in at the borders.' An apparent imperfection, but in reality an advantage to concentration of attention, since by the rapidity and exactness with which the eye can be turned from point to point, we can at will focus and minutely examine any spot; any such spot becomes the centre of the picture, and a focus of attention, as well as a focus of vision.

The extent of the field of vision in any given case is measured by the *perimeter*; this is in principle a hemisphere, at the centre of which the eye under examination is placed (the other eye being closed), while the examiner determines on the hollow surface of the hemisphere those points at which the patient just ceases or just commences to see a small object moved in various directions. The observations plotted on skeleton charts, graduated in degrees, give in graphic form the field of vision of the stationary eye.

*Scheiner's experiment* is deserving of careful repetition by any one who desires to realise from a different point of view facts relating to

accommodation, and in addition to this the formation of double images in a single eye, and the relation of retinal impressions to cerebral perceptions.

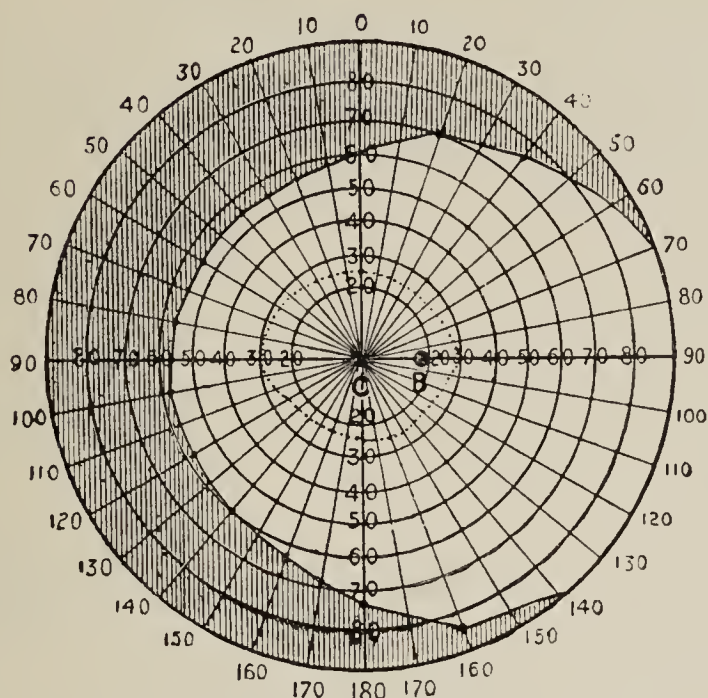


FIG. 232.—PERIMETRIC CHART OF THE RIGHT EYE.

The thick line surrounds the area within which *white* is visible, the point C being fixed by the eye (right); the fine dotted line surrounds the area within which green is visible. The area of red would be somewhat larger, that of blue larger still, though not so large as that of white. B = the situation of the blind spot.

Two pinholes are pricked in a card at about one millimeter from each other; and the card being held in front of one eye, with the two pinholes side by side in front of the pupil, a needle, held vertically, is looked at through them.

(a) With the eye accommodated for the needle, the latter is seen single, and if it be brought gradually nearer to the eye, a point is

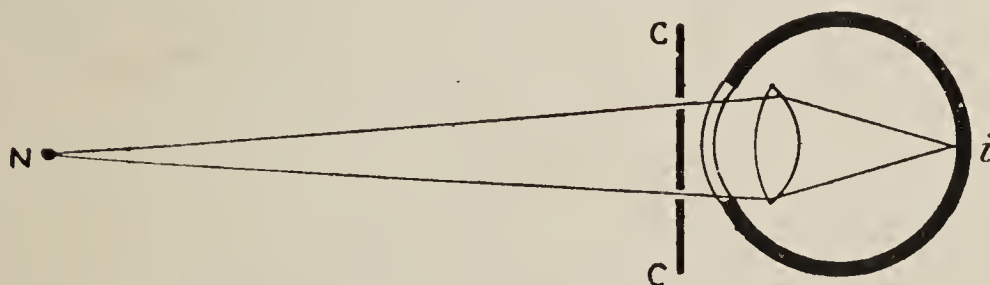


FIG. 233.—TO ILLUSTRATE SCHEINER'S EXPERIMENT. See text (a).

Vertical section of eyeball. N, section of needle held horizontally; C C, card pierced by two pinholes, one above the other, held vertically;

Or, horizontal section of eyeball. N, section of needle held vertically; C C, card held vertically, with pinholes right and left.

reached at which it can no longer be seen single by any effort of accommodation; the distance of the needle in front of the eye is now that of the near point of distinct vision. With the eye accommodated to a point either nearer or farther than the needle, the latter is seen double.

(b) If the eye is accommodated to a point beyond the needle, or, better still, if it is non-accommodated, two images will appear, the eye



being under-accommodated as regards the needle; these images will move towards each other as the needle is moved from the eye, will

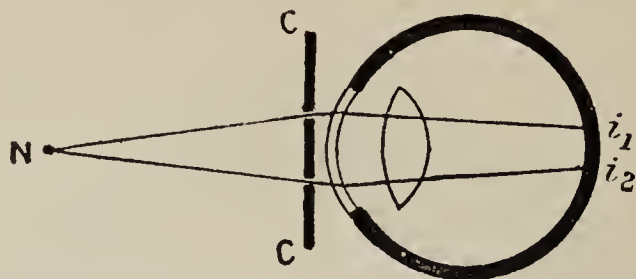


FIG. 234. - TO ILLUSTRATE SCHEINER'S EXPERIMENT. See text (b).

Vertical section of eyeball and horizontal needle, *or* horizontal section of eyeball and vertical needle.

Blocking the upper hole in the card abolishes the retinal image  $i_1$ , *i.e.* the apparent inferior needle in the field of vision; blocking the lower hole abolishes  $i_2$ , *i.e.* the apparent superior needle.

enlarge and move away from each other as the needle is moved towards the eye. If the right-hand hole is covered, the apparent left-hand needle will disappear; and, *vice versa*, if the left-hand hole is blocked the apparent right-hand needle will be lost.

(c) If the eye is accommodated to a point nearer than the needle, it is over-accommodated as regards the needle; two images will be apparent, and if the needle be moved to and from the eye, they will move to and from each other. Blocking the right-hand hole effaces the apparent right-hand needle, blocking the left-hand hole effaces the apparent left-hand needle.

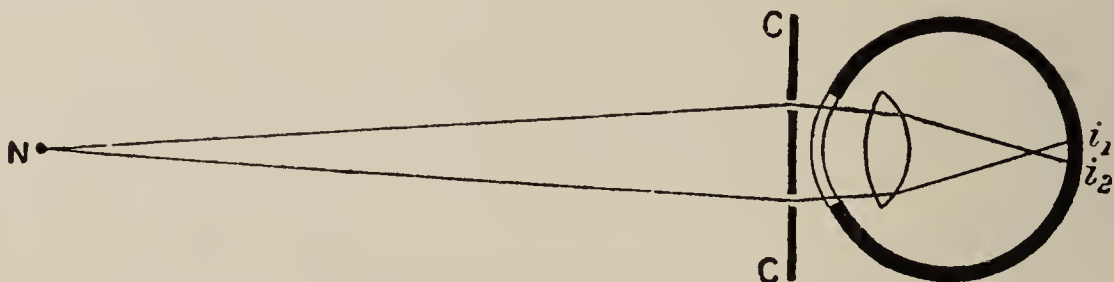


FIG. 235.—TO ILLUSTRATE SCHEINER'S EXPERIMENT. See text (c).

Vertical section of eyeball and horizontal needle, *or* horizontal section of eyeball and vertical needle.

Blocking the upper hole abolishes the retinal image  $i_2$ , *i.e.* the apparent superior needle; blocking the lower hole abolishes  $i_1$ , *i.e.* the apparent inferior needle.

The whole series of observations may be repeated with the needle held horizontally, the card being held so that the holes are one above the other in front of the pupil.

*Chromatic dispersion.*—The lens of the eye, like other lenses, refracts blue rays more strongly than red rays. Consequently the focus of blue is nearer to the lens than the focus of red. A spot of white light is surrounded by a blue halo if the eye is focussed to a point beyond the spot, by a red halo if it is focussed to a point nearer than the spot; to see these halos distinctly the light should be viewed through cobalt glass, so as to cut off the yellow and green, allowing only red and blue to pass.

A blue light and a red light at the same distance from the eye appear to be unequally distant; the red light, requiring greater accommodation than the blue, appears to be the nearer of the two, nor can both lights be exactly focussed together upon the retina. Of a red and a blue surface side by side, the red appears to be the nearer.<sup>1</sup> The difference is such that a normal eye focussed for parallel red rays is at the same time focussed for blue rays divergent from the points of a surface about 3 feet distant.

*Astigmatism.*—The surface of the cornea is not perfectly spherical; the curvature of its vertical meridian is usually greater than that of its horizontal meridian, and the difference may be so pronounced as to interfere with the distinct focussing of objects. With a cornea of such curvatures ('spoon-shaped'), a point of light cannot form a focal point upon the retina (hence the name astigmatism), but forms a linear focus; if the eye is adjusted so that the rays diverging from the point in the vertical plane (meridian of greater curvature) come to a

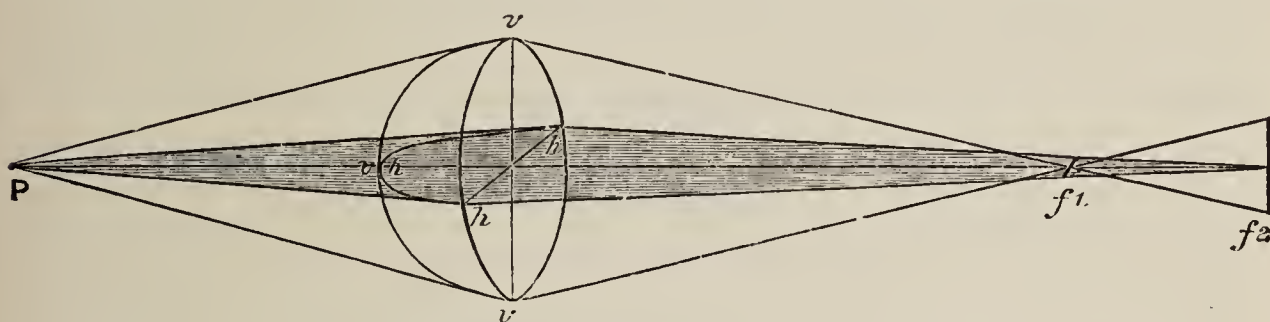


FIG. 236.—REGULAR ASTIGMATISM.

The curvature of the cornea is greater in the vertical meridian,  $v v v$ , than in the horizontal meridian,  $h h h$ . The point of light, P, consequently has a first linear focus at  $f_1$  which is horizontal, and a second linear focus at  $f_2$ , which is vertical. If, instead of P, the object were a vertical and a horizontal line crossed, the vertical line would be in focus at  $f_2$ , the horizontal line would be in focus at  $f_1$ . To bring the two lines into focus at the same time on the same plane it would be necessary to use a convex cylindrical glass to add to the horizontal convexity  $h h h$ , or a concave cylindrical glass to subtract from the vertical convexity  $v v v$ .

focus on the retina, then the rays diverging from the point in the horizontal plane (meridian of lesser curvature) will reach the retina before they have come to a focus (*i.e.* be under-focussed), and a horizontal linear focus will be formed; *vice versa*, if the eye is adjusted so that rays in the horizontal plane (meridian of lesser curvature) are focussed upon the retina, then the rays in the vertical plane will reach the retina after coming to a focus (*i.e.* be over-focussed), and a vertical linear focus will be formed. As shown in the diagram, the horizontal linear focus,  $f_1$ , is nearer than the vertical linear focus,  $f_2$ .

Consider next the case of a series of points, viz. a line, vertical or horizontal. A person whose cornea is regularly astigmatic, as supposed above, can see distinctly either a horizontal or a vertical line, but not

<sup>1</sup> This is not a complete explanation. Einthoven has shown that the effect is mainly dependent upon excentricity of the pupil. Most persons see red in front of blue, but some persons see blue in front of red.





brought close to the eye, and a retinal vessel in the plane  $f$  will have its real reversed image nearer to the eye in the plane  $f'$ . An observing eye placed at  $o$  (so that the distance  $o f'$  is greater than that of the observer's near point) could see this image of the retinal vessels if the

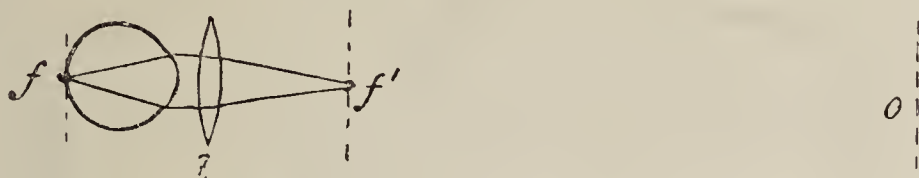


FIG. 238.

retina were a source of light or lighted up so as to reflect sufficient light. This can be done. The fundus of the eye is made visible by rays reflected from a mirror, which light up its vessels, &c.; rays reflected from parts so illuminated emerge from the observed eye and form the image  $f'$ , which is seen by the observing eye.

The main principle upon which the ophthalmoscope depends is thus that the line of vision of the observing eye shall be in the line of illumination. This is effected by means of the perforated mirror through which the observer looks while he directs light along his line of vision.

This method of examination is known as the *indirect method*. The retinal object,  $a b$ , is illuminated by light reflected from a concave perforated mirror; rays reflected from the bright object,  $a b$ , are refracted

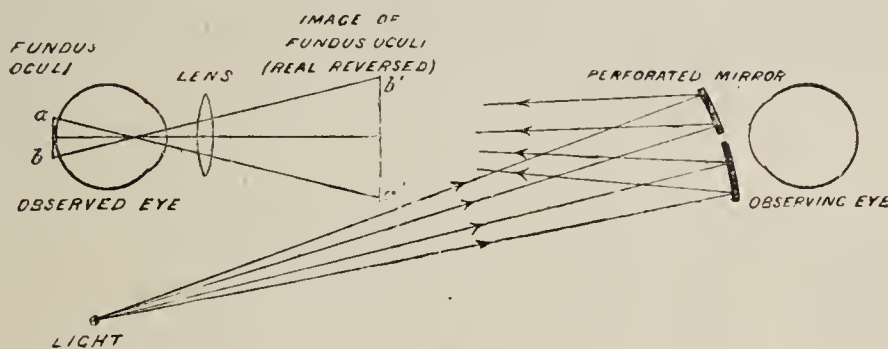


FIG. 239.—DIAGRAM TO SHOW HOW OBJECTS ON THE FUNDUS OCULI ARE ILLUMINATED AND SEEN BY THE INDIRECT METHOD.

The lines of vision and of illumination are made to coincide by the perforated mirror, which is concave in order to concentrate the light.

by the observed eye and auxiliary lens to form a real reversed image,  $b' a'$ , which is looked at by the observing eye through a hole in the mirror (fig. 239).

The second method of examination is known as the *direct method*; the image thus obtained is a virtual erect image formed by rays as they emerge from the observed eye. In this case, to obtain any useful view, the observing and observed eyes must be as close as possible. As will be understood from the description of the three conditions of refraction that we may have to deal with, the 'direct method' is one of the means of detecting and estimating errors of refraction.

Rays emerging from any given point at the fundus of a normal eye



will be parallel, from a short eye they will be divergent, from a long eye they will be convergent. Such rays will not come to a focus in the

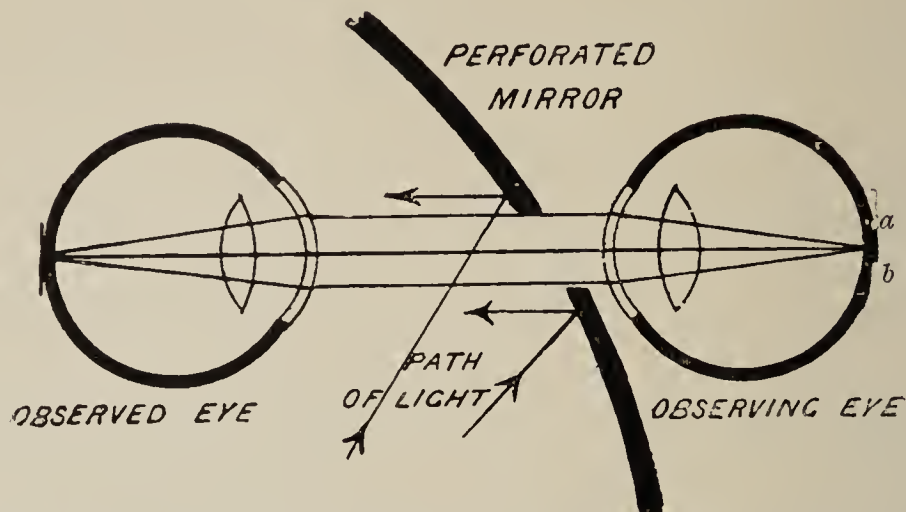


FIG. 240.—DIAGRAM TO SHOW HOW OBJECTS ON THE FUNDUS OCULI ARE ILLUMINATED AND SEEN BY THE DIRECT METHOD.

Both the eyes are supposed to be emmetropic and non-accommodated; under these conditions the vessels visible in the observed eye are slightly anterior to the rod-and-cone layer, which is supposed to be in focus, and rays from the vessels would be slightly divergent. The size of the mirror-aperture is greatly exaggerated.

case of the normal eye (so long as its accommodation is entirely relaxed); they will have a virtual focus in the case of the short eye, a real focus in the case of the long eye; in the first case (E) no image of the retinal surface is formed, in the second (H) a virtual erect image is formed behind the eye, in the third (M) a real reversed image is formed in front of the eye (fig. 241).

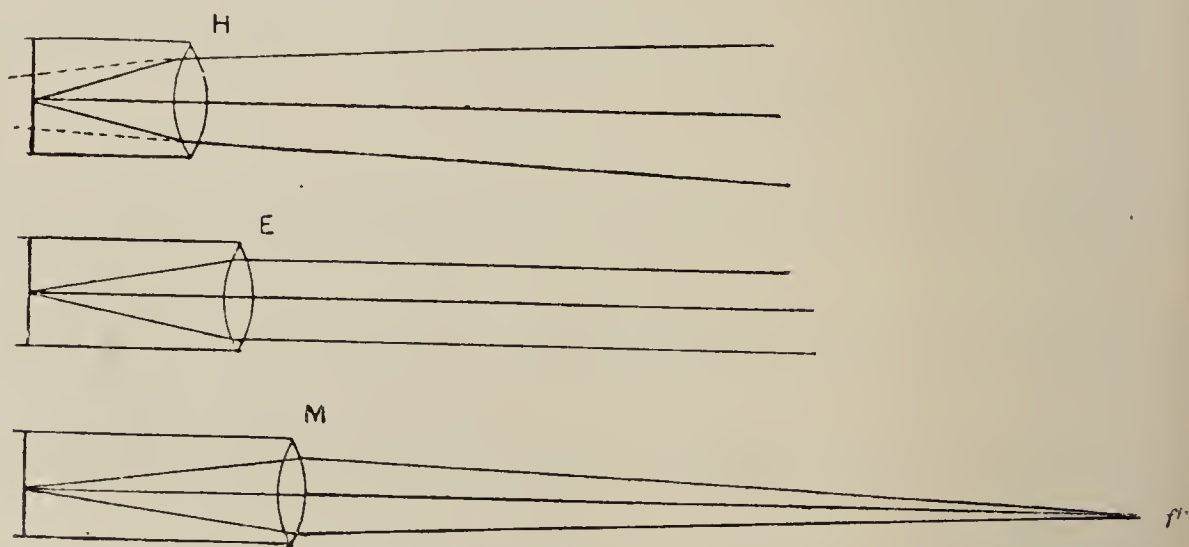


FIG. 241.—PATH OF RAY EMERGING FROM A HYPERMETROPIC, FROM AN EMMETROPIC, AND FROM A MYOPIC EYE.

From the emmetropic eye, E, with relaxed accommodation, the rays will be parallel and give no image.

From the hypermetropic eye, H, the rays will be divergent, and, being prolonged backwards, give a virtual erect image.

From the myopic eye, M (or from an accommodated emmetropic eye), the rays will be convergent and give a real reversed image at  $f'$ .

The chief practical difficulty in the use of the direct method is for

the observer to maintain his eye non-accommodated. Thus, to estimate hypermetropia he has to find the strength of the convex lens which, added to his non-accommodated eye, gives distinct vision of the fundus, and it is therefore essential that he should not unconsciously exert any effort of accommodation. For the minute examination of a normal eye by the direct method the observer's eye must be non-accommodated, for a myopic eye it must be less than non-accommodated, *i.e.* the assistance of a concave lens is required; on the other hand, for a hypermetropic eye the observer has to accommodate or to use a convex lens behind the mirror.

The observed eye also must be non-accommodated in order that its refraction may be estimated, as well as for the minute examination of the fundus. Whereas rays emerging from a normal non-accommodated eye are parallel, rays emerging from a normal accommodated eye will be converging, *i.e.* such an eye will be equivalent to the myopic eye, and its vessels will not be seen by close examination unless a concave lens is used behind the mirror. Hence it will be understood that in the direct examination of the fundus: (1) a concave lens must be added behind the mirror if the observed eye is myopic; (2) a weak concave lens may be added if the observed eye is normal, in which case the observer accommodates the fundus; (3) a convex lens may be used if the observed eye is extremely hypermetropic.

**Relation of retina to field of vision.**—In performing Scheiner's experiment the observer will have noticed that by blotting out the image which is superior, inferior, right or left as regards its retinal position, he loses sight of the apparent image in his field of vision which is inferior, superior, left or right. In other words, impressions made on the upper, lower, right or left part of the retina excite in consciousness the sensations that objects are situated below or above, to the left or to the right, of the line of vision. We may easily recognise this to be a necessary relation when we have realised that the retinal image is reversed, objects at our feet being focussed on the upper portion of the retina, objects above our heads on its lower portion, objects to our left on its right half, objects to our right on its left half. If, as sometimes occurs, the lateral halves of the two retinae are paralysed, say on the right side (right hemiopia), the left half of the field of vision is blotted out (left hemianopia); *vice versa*, left hemiopia of

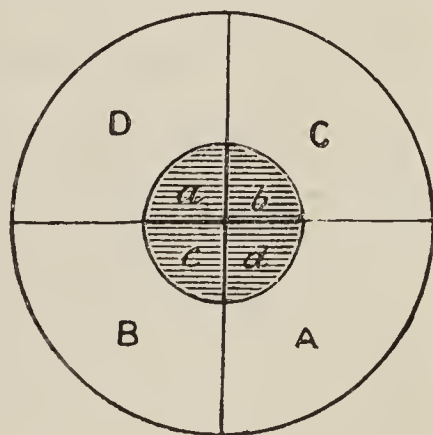


FIG. 242.

The quadrants A B C D of the field of vision are projected upon the quadrants a b c d of the retina.



the two retinæ will cause right hemianopia as regards the field of vision.<sup>1</sup>

Moreover, we normally see objects single with both eyes open, although two retinal images are formed, one in each eye. This is

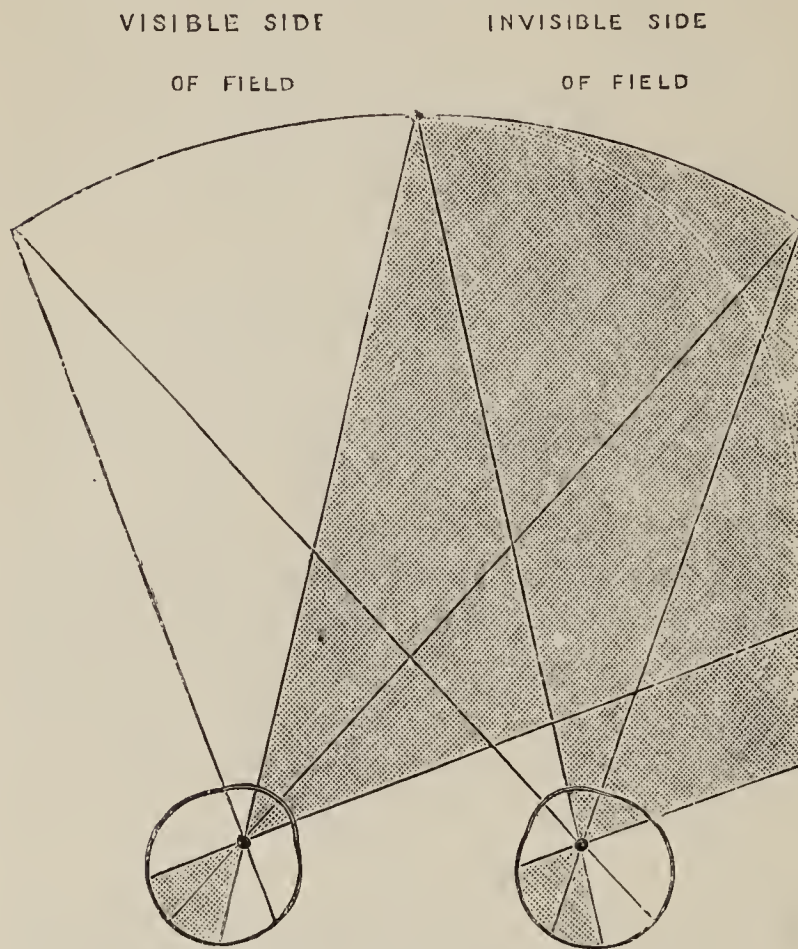


FIG. 243.—LEFT HEMIOPIA AND RIGHT HEMIANOPIA.

due to the fact that when an object is looked at, its image (and those of its immediate surroundings) are focussed upon *corresponding parts* of the two retinæ. The doctrine of corresponding points will, however, be best understood by taking into account the circumstances under which, correspondence not being effected, double vision occurs. Before so doing it should be remarked that in

Scheiner's experiment we have experienced double vision with only one eye open; this obviously being due to the fact that we have, by means of the two pinholes, made two images of the same object fall upon different parts of the same retina.

*Corresponding points.*—Single vision with the two eyes and double vision with the two eyes differ in this respect, that in the former the object gives on each side of the two retinæ an image, every point of which comes to a focus upon each of two corresponding points in the two retinæ, while in the latter the two retinal images are not outlined upon corresponding points. In the first case the excitations caused by the two corresponding images fuse in consciousness and cause perception of a single

<sup>1</sup> The unguarded use of these terms in various senses leads to much unnecessary confusion, and their meaning is sometimes only to be gathered from the context. Hemioopia is literally half-vision, while hemianopia (or hemianopsia) is half-blindness. The expression 'left hemioopia,' as used by different writers, may mean paralysis of the right or left halves of the retina, or blindness to the left or to the right of the subject. Ambiguity is avoided by stating the sides to which vision is preserved or lost. See also p. 543.

object ; in the second, the excitations of the two non-corresponding images do not so fuse, and the perception of an apparently double object is their consequence. Corresponding points are thus physiological pairs of points, one in the right eye, one in the left eye, from the simultaneous stimulation of which a single optical sensation results ; anatomically, they are almost exactly the symmetrical points in the two retinae that would cover each other in pairs if the two retinae were superposed one upon the other, with the yellow spots as centre.

The yellow spots correspond in the two eyes, and when an object is looked at, its image is

formed upon each of the two yellow spots. The blind spots in the two eyes do not correspond ; being situated in the nasal segment in each eye, they would not cover each other if the retinae were superposed as above described. The nasal side of the left eye corresponds with the temporal side of the right eye, and the temporal side of the left eye with the nasal side of the right eye.

*The horopter.*—The horopter is that surface or series of points in space the images of which fall upon corresponding points of the two retinae. A complete exposition of the form assumed by the horopter with various positions of the eyeball would necessitate a mathematical analysis far beyond the scope of a simple text-book of physiology ; we shall here restrict ourselves to the verbal definition of the ‘horopter,’ and to the geometrical definition of Müller’s horopter circle, which occurs when the visual axes converge upon a near object.

In this position the horopter is formed by a horizontal circle passing through the object, and through the nodal points of the two eyeballs, and by a straight line drawn through the point of fixation in the median plane tilted away from the observer. Müller’s horopter circle is a practical illustration of prop. 21 of Book III. of Euclid, in which it is demonstrated that any given chord of a circle subtends equal angles at all points of the circumference, or, otherwise expressed, that the *locus* of the vertices of all triangles on the same side of the same base, with a constant angle at the vertex, is the arc of a circle.

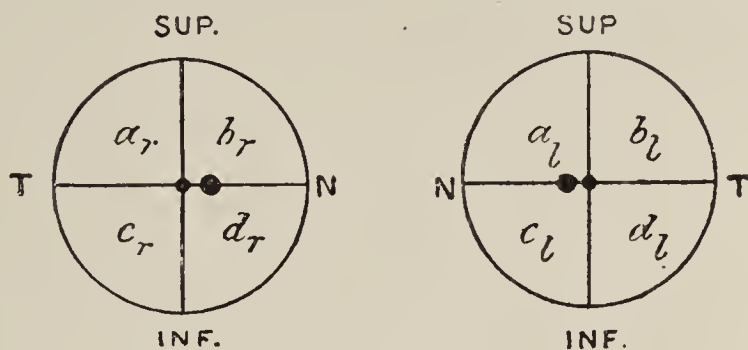


FIG. 244.

$a_r, b_r, c_r, d_r$ , and  $a_l, b_l, c_l, d_l$ , are corresponding quadrants in the two retinae, and would exactly cover each other if superposed. Each quadrant may similarly be imagined to be symmetrically divided into symmetrical corresponding parts, and each such part further subdivided into single rods and cones symmetrically situated in the two retinae, *i.e.* corresponding. The blind spots are non-corresponding.



In the primary position, with the visual lines parallel and the visual plane horizontal, the horopter is formed by the distant vertical view if the retinal meridians are at absolute right angles to each other, or by a horizontal surface below the feet if—as is usually the case—the vertical retinal meridian is not at right angles to the horizontal meridian; but the construction of the horopter in these cases would transgress our limits.<sup>1</sup>

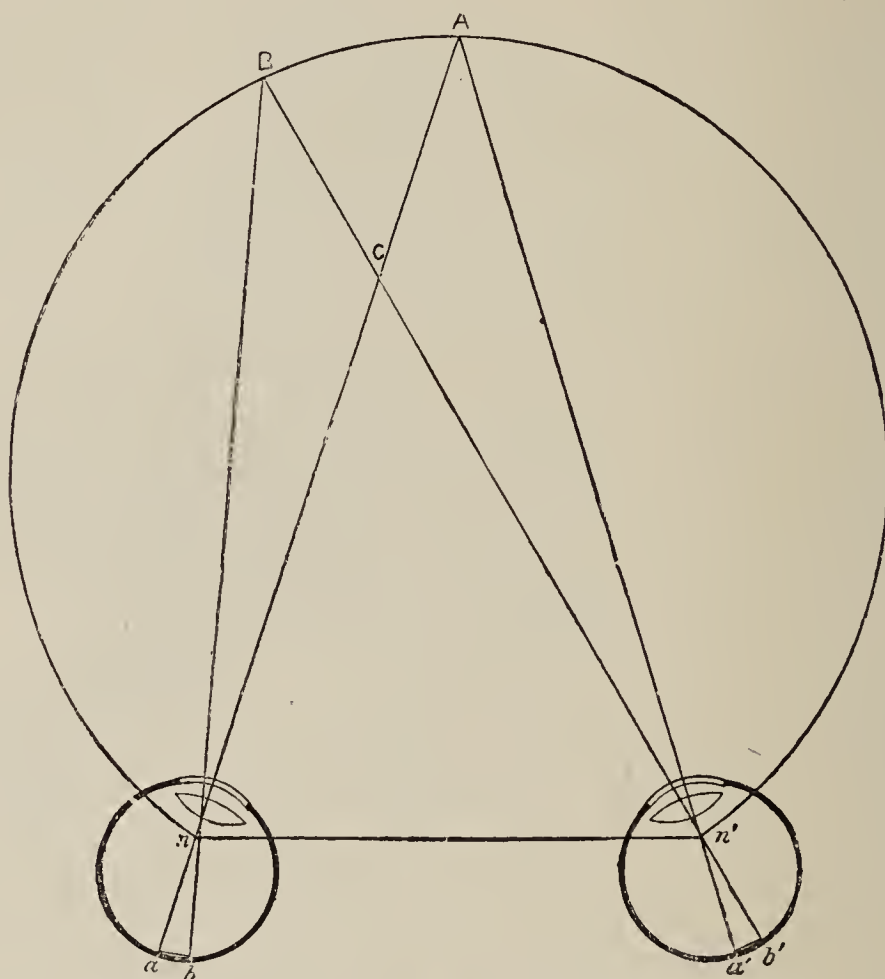


FIG. 245.—TO ILLUSTRATE MÜLLER'S HOROPTERIC CIRCLE.

A is the point of regard; its retinal images are formed on the yellow spots at  $a, a'$ .  $b, b'$  are corresponding points of the two retinae, therefore the distance  $ab =$  the distance  $a'b'$ .  $n, n'$  are the nodal points of the two eyes. The angles  $anb, a'n'b'$ , are equal to each other and to the opposite angles,  $AnB, A'n'B$ ; and in the triangles  $BCn, ACn'$ , the opposite angles at C are equal; therefore, in these triangles the remaining angles at A and B are equal. Therefore the point B by which the corresponding images  $b, b'$  are formed must lie in the arc of a circle passing through the points B,  $n$ , and  $n'$ . Similarly, it may be shown that any other point forming corresponding images on the two retinae will lie in the circular arc  $nAn'$ .

*Double vision.*—Double vision of an object with both eyes open is most apparent when the gaze is directed to a point nearer or farther than the object, in or nearly in the same line of vision. If a finger is held up in front of the face at arm's length while the eyes are adjusted to a more distant point, the finger will be

<sup>1</sup> Students desirous of further information on the subject may refer to Helmholtz, *Physiological Optics*, p. 745; Hering, *Beiträge z. Physiol.* 1864; and in Hermann's *Handbuch d. Physiol.* vol. iv.

seen double; closure of the left eye will now abolish the finger apparently seen to the right, and *vice versa*. If two fingers are held up in front of the face, one at arm's length, the other at half-arm's length, while the eyes are adjusted to distinct vision of the nearer finger, the more distant finger will be seen double; closure of the left eye will now abolish the finger apparently seen to the left, and *vice versa*.

*Double vision from squint (Diplopia).*—Double vision also occurs when *both* visual axes are not directed towards an object that is being looked at, *i.e.* when there is squint or *strabismus* of one eye. The subject of squint, when looking at an object, directs to it the visual axis of the sound or working eye, while that of the squinting eye diverges from it, usually to the right or to the left. Consequently the images on the two retinae are not formed upon corresponding parts; that formed upon the working eye is called the true image, that upon the squinting eye is called the false image. The position of the 'false' in relation to the true image in various cases of misdirection of a visual axis, will be best understood when the normal movements have been considered.

#### Movements of the eyeball. —

As has been stated above, it is a necessary condition to single vision with the two eyes that the objects should form their images upon corresponding parts of each retina—upon the yellow spots when objects are looked at. Now single vision is the normal event, and is preserved with all kinds of direction of the lines of vision; its necessary condition

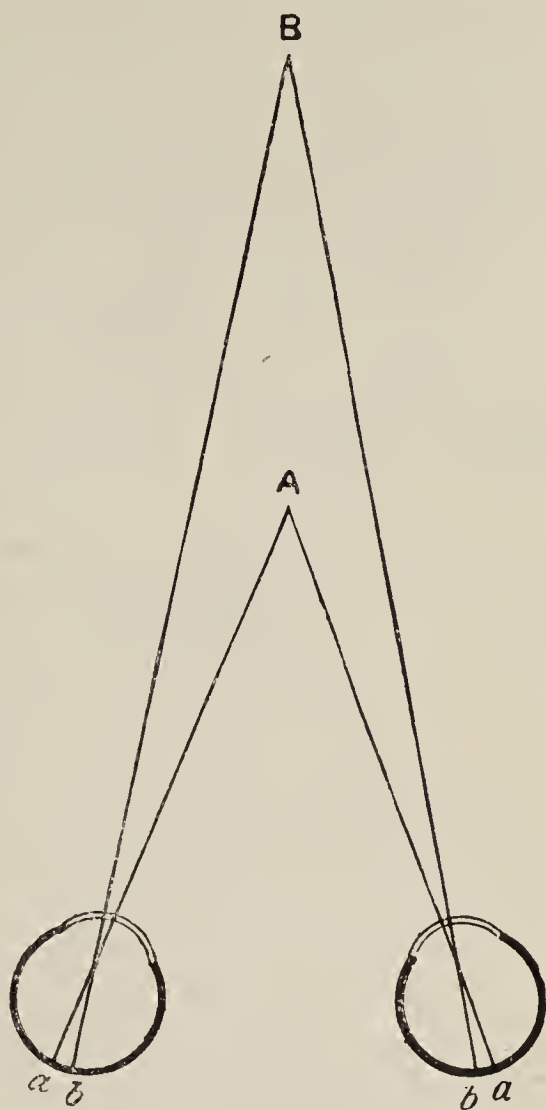


FIG. 246.

*Double vision of an object for which the eyes are not accommodated.*

The eyes are focussed on B, *i.e.* on the yellow spots at *b, b*.

∴ A has retinal images at *a, a*.

On closing the left eye, the apparent right finger, A, vanishes.

On closing the right eye, the apparent left finger, A, vanishes.

The eyes are focussed on A, *i.e.* the yellow spots are at *a, a*.

∴ B has retinal images at *b, b*.

On closing the left eye the apparent left finger, B, vanishes.

On closing the right eye the apparent right finger, B, vanishes.



is, then, a very delicate co-ordination of the muscles that move the eyes. Each eye is moved by six muscles: four recti—superior, inferior, internal, and external; two obliqui—superior and inferior. The third, or motor oculi nerve supplies the superior, inferior, and internal recti, and the inferior oblique; the fourth, or trochlearis nerve supplies the superior oblique; the sixth, or abducens nerve supplies the external rectus. The directions in which the eyeball is rotated are best defined by referring to the direction in which

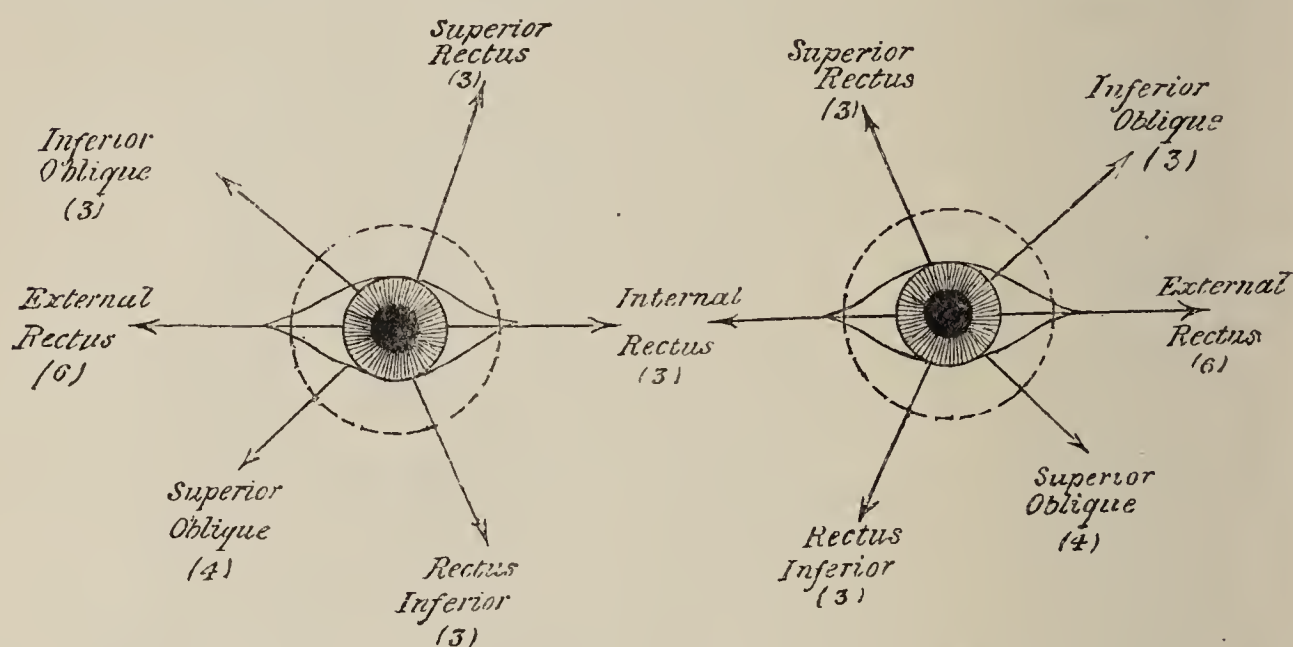


FIG. 247.

Diagram to illustrate the directions towards which the pupil is moved by the separate action of the six muscles of the eyeball. The eyes are turned inwards and outwards by the external and internal recti; the internal rectus of one side is the yoke-fellow of the external rectus of the opposite side in these conjugate movements. The eyes are turned upwards by the superior rectus and inferior oblique, downwards by the inferior rectus and superior oblique.

the cornea is moved, and the effects of the individual muscles in this respect are expressed in the diagram by arrows indicative of the direction towards which the cornea moves.

For the six muscles of either eye it is evident that

the internal rectus rotates the eye *inwards*

„ external rectus	„ „	<i>outwards</i>
„ superior rectus	„ „	<i>upwards and inwards</i>
„ inferior rectus	„ „	<i>downwards and inwards</i>
„ superior oblique	„ „	<i>downwards and outwards</i>
„ inferior oblique	„ „	<i>upwards and outwards</i>

and that rotation vertically *upwards* is effected by the joint action of the superior rectus and of the inferior oblique, rotation vertically *downwards* by the joint action of the inferior rectus and of the superior oblique. It is also evident from the diagram that antagonistic pairs of muscles are:—internal rectus and external

rectus; superior rectus and superior oblique; inferior rectus and inferior oblique.

The eyeball is thus, as regards its principal movements, a sphere capable of rotating in any direction round its centre, up, down, in and out; vertical movements being effected round a horizontal axis, horizontal movements round a vertical axis, oblique movements round oblique axes, there being under ordinary circumstances little or no torsion of the globe round its antero-posterior axis. The various axes round which the eye rotates all lie in one plane, at right angles to the line of vision, and passing through the centre of rotation, the exact position of which is 1.77 mm. behind the centre of the eyeball. This plane is known as Listing's plane. With the exception of the vertical axis, round which the eyeball is rotated by the internal and external recti, the axes of rotation of single muscles do not coincide with the axes of physiological movements. The latter, as just stated, lie in Listing's plane, which is approximately vertical; the former lie in a horizontal plane, as shown by fig. 248—the recti superior and inferior, directed forwards and outwards, would respectively rotate the eye up and down round the axis *rr*; the obliqui, directed from before backwards and outwards, would rotate the eye round the axis *oo*. But they never normally do so; the ordinary movements of the eyeball are rotations effected by the simultaneous action of several muscles.

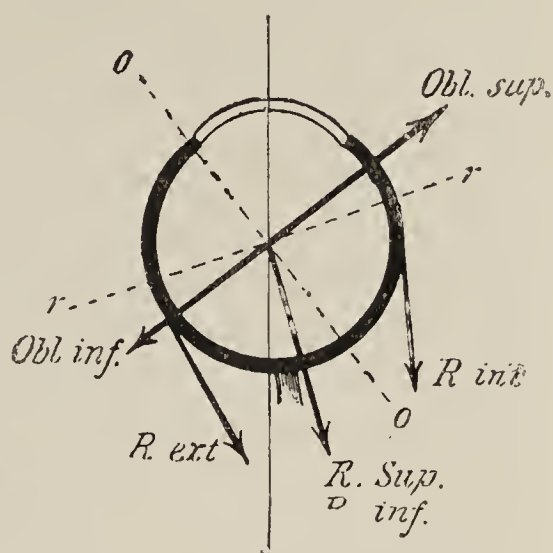


FIG. 248.—HORIZONTAL SECTION OF LEFT EYEBALL.




To illustrate the direction of action and the axes of rotation of its muscles. The arrows indicate the direction of action, the dotted lines indicate axes of rotation. *oo* is the axis round which the eyeball rotates by action of the obliqui. *rr* is the axis of the superior and inferior rectus. The axis of the superior and inferior rectus would be represented by a pin through the point of intersection of *oo* and *rr*, perpendicular to the plane of the paper.

*Wheel-movements of the iris—true and false.*—In these parallel movements there is no rotation of the eyeball round an antero-posterior axis (wheel-movement of iris); the wheel-movement or torsion apparently indicated by the following experiment is not real, but a mixed effect, due in part to an error of judgment, in part to inclinations that a horizontal retinal meridian suffers in oblique movements round oblique transverse axes. It is convenient to refer these effects to



‘wheel-movements,’ but we must then characterise them as ‘false wheel-movements’ in distinction from ‘true wheel-movements,’ which do actually take place under certain conditions, viz. with converging visual axes, and with inclinations of the head towards either shoulder while the visual axes are kept fixed; in the first case the two eyeballs rotate *outwards* round the axes of the lines of vision, in the second case the two eyeballs rotate *against* the direction towards which the head is inclined.

The apparent rotation, or false wheel-movement, is demonstrated by means of after-images, as follows:—The observer sits with the gaze fixed in the primary position (*i.e.* head erect—eyes looking horizontally straight forward), opposite to but not too near a grey wall with faint vertical lines; a strip of red paper is fixed vertically against the wall, on a level with the eyes. The centre of the strip having been steadily regarded for a few seconds, the eyes are suddenly directed vertically up or down, or horizontally to the right or left. The green-blue after-image of the strip will in all these cases remain vertical; but if the eyes are directed obliquely upwards and to the right, or downwards and to the left, the after-image will appear tilted to the right; if the eyes are directed obliquely upwards and to the left, or downwards and to the right, the after-image will appear tilted to the left.

That these rotations are *apparent* is at once recognised by substituting a horizontal for a vertical red line, ‘branding’ the retina as before with a green-blue after-image; the latter will be inclined *against* the inclinations of the vertical after-images. Or, if a rectangular cross be taken, the vertical and horizontal components of the after-images will set towards each other. The explanation is as follows:—A rectangular cross  seen upwards and to the right casts a retinal image that is not rectangular but inclined thus , yet it is judged to be rectangular; the actually rectangular brand projected upwards and to the right is therefore misjudged, and interpreted as if due to a cross inclined thus ; and similarly for the three other cases—down to right, down to left, up to left. The best way to avoid this complication by misjudgment is to take a single coloured line and a faint lined surface, both movable round a central axis, and to adjust the lines obliquely, in the directions along which the line of vision is to be shifted from the first point of fixation.

From a knowledge of the action of muscles on the eyeball and of the relation between retinal images and apparent position of objects in the field of vision, it is easy to deduce what must be the displacement of the eyeball and the position of false images in the field of vision in consequence of paralysis of one or more muscles or nerves.

A simple case is that of the *external rectus* ; its action is to rotate the eye outwards, the right eye to the right, the left eye to the left ; if it is paralysed the eye will squint inwards, the right eye to the left, the left eye to the right ; in consequence of this displacement the image of an object looked at by the patient will fall upon the retina on the internal or nasal side of the fovea, and the apparent image in his field of vision will be on the external or temporal side of the true image ; if the right external rectus is paralysed the false image (seen by means of the right eye) will lie on the right side of the true image (seen by the left eye) ; if the left external rectus is paralysed the false image (by the left eye) will lie on the left side of the true image (by the right eye). In neither case will the false image be tilted, but simply displaced laterally.

Name		Normal action and direction in which movements are defective in paralysis	Direction of paralytic squint caused by excessive action of antagonists	Relation of false image on retina to position it should occupy	Relation of apparent image in field of vision to position of object seen by working eye	Inclination of apparent image to or from working image
Internal rectus . . .	3	In	Out	Ext.	In	None
<b>External rectus</b> . . .	6	Out	In	Int.	Out	None
Superior rectus . . .	3	Up and in	Down and out	Inf. and ext.	Up and in	From
Inferior oblique . . .	3	Up and out	Down and in	Inf. and int.	Up and out	From
Inferior rectus . . .	3	Down and in	Up and out	Sup. and ext.	Down and in	To
<b>Superior oblique</b> . . .	4	Down and out	Up and in	Sup. and int.	Down and out	To
<b>Third nerve</b> . . .		Up and in	Down and out	Inf. and ext.	Up and in	From
<hr/>						
Internal rectus . . .	R	L	R	R	L	None
" " " " " "	L	R	L	L	R	None
<b>"External" rectus</b> . . .	R	R	L	L	R	None
<b>or Sixth Nerve</b> . . .	L	L	R	R	L	None
Superior rectus . . .	R	Up and L	Down and R	Inf. and R	Sup. and L	To L
" " " " " "	L	Up and R	Down and L	Inf. and L	Sup. and R	To R
Inferior oblique . . .	R	Up and R	Down and L	Inf. and L	Sup. and R	To R
" " " " " "	L	Up and L	Down and R	Inf. and R	Sup. and L	To L
Inferior rectus . . .	R	Down and L	Up and R	Sup. and R	Inf. and L	To R
" " " " " "	L	Down and R	Up and L	Sup. and L	Inf. and R	To L
<b>"Superior" oblique</b> . . .	R	Down and R	Up and L	Sup. and L	Inf. and R	To L
<b>or Fourth Nerve</b> . . .	L	Down and L	Up and R	Sup. and R	Inf. and L	To R
<b>Third nerve</b> . . .	R	Up and L	Down and R	Inf. and R	Sup. and L	To L
" " " " " "	L	Up and R	Down and L	Inf. and L	Sup. and R	To R

A more complex case is that of the *third nerve* : its action is to rotate the eye upwards and inwards ; if the third right nerve is paralysed, the right eyeball squints downwards and to the right ; the false image on the retina of that eye will be below and to the right of the yellow spot ; the apparent image in the field of vision will be, in relation to the true image, above and to the left. If the third left nerve is paralysed, the left eyeball squints downwards and to the left : the false image on the retina will be below and to the left of the yellow spot ; the apparent image in



the field of vision will be, in relation to the true image, above and to the right.

Reasoning in a similar manner from the known action of a given muscle to the displacement which the eyeball will theoretically undergo in consequence, it is easy to construct a

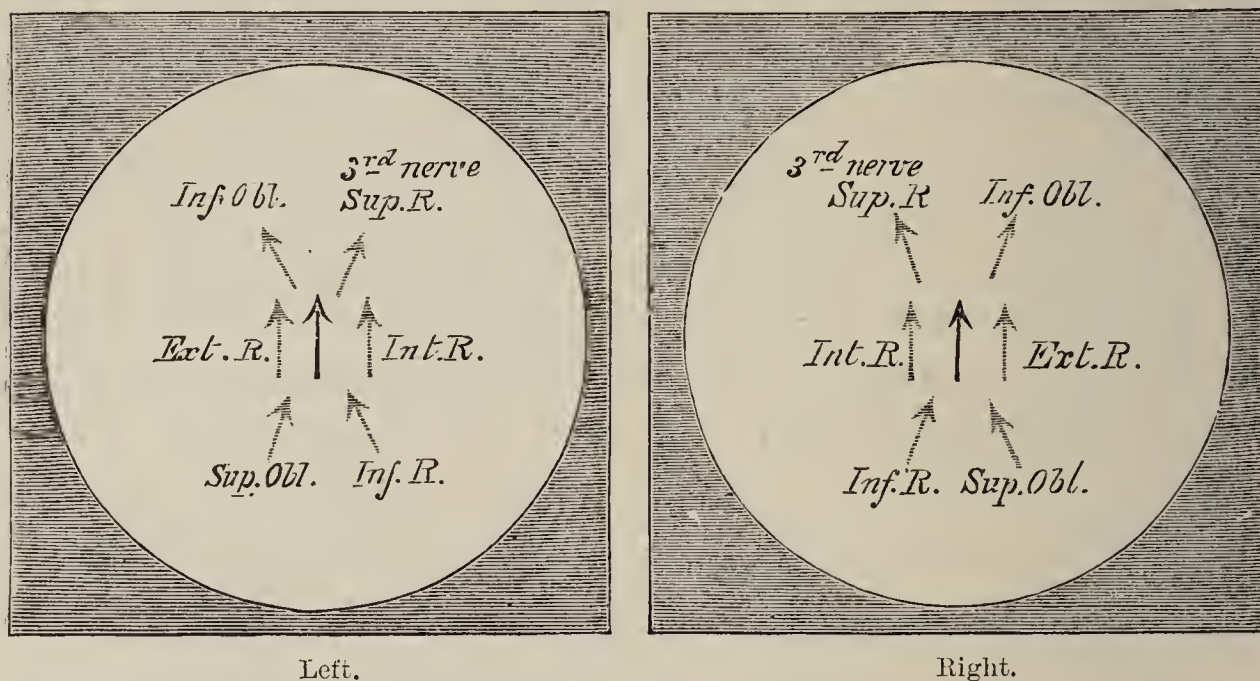


FIG. 249.—APPARENT DISPLACEMENTS OF AN OBJECT IN THE FIELD OF VISION IN CONSEQUENCE OF PARALYSIS OF THE SEVERAL MUSCLES OF THE EYEBALL.

The central arrow is the position of the object in the field as viewed by the working eye. N.B.—From the statements in the text it will be evident that fig. 249 may be constructed from fig. 247. The latter viewed by transparency or in a mirror gives the construction of the former. It will thus be realised that (1) the displacement in consequence of the paralysis of any muscle is the converse of its displacement by the action of that muscle; (2) the accompanying displacement of the false image in relation to the true image, in consequence of paralysis of a muscle, is in the same direction as the normal action of that muscle.

These rules apply equally to the resultant effects of any nerve and of any group of muscles. Practically, however, the great majority of cases that actually present themselves are those which are indicated by heavy type in the table (p. 445), and in connection with these common cases the effects are usually complicated by collateral effects, the description of which belongs to ophthalmology. One of these collateral effects is of physiological import, and may therefore be alluded to here. In normal conjugate movements of the two eyes, *e.g.* to the right or left, the external rectus of one eye acts with the internal rectus of the opposite eye (*vide* p. 533); if, for instance the right external rectus is paralysed, the right eye squints inwards and cannot be directed to the left, but the attempt to do so causes an excessive action of its yoke-fellow, the left internal rectus, and the left or normal eye now squints inwards. This is called a secondary squint, and has been compared to the effect that would be produced in driving a pair of horses with unequally sensitive mouths—an effort to turn the pair would cause the normal horse (or normal eye) to deviate much more than the hard-mouthed horse (or paralysed eye).

table of all the different possible cases, and to embody them in a diagram (fig. 294 constructed from the diagram representing the movements of the eyeballs (fig. 247).

We have described movements of the eyeball, for the sake of simplicity, without reference to the action of antagonist muscles.

It is probable, however, that such muscles *contract* simultaneously with the muscles directly producing deflections of the line of vision; this check action contributes to that fixation of the eyeball without which a steady sighting would be impossible. For instance, when the eyes are turned to the right, the right external rectus is checked by the right internal rectus, and the left internal rectus is checked by the left external rectus; in converging the visual axes upon a near object, the two internal recti are checked by the two external recti. Close reading or sewing necessitates strong convergence and strong accommodation; an excessive pressure from this cause during adolescence can eventually lengthen the eye and produce myopia, as was once illustrated on a large scale in German schools.

But it may also happen in the synergic action of the ocular muscles that an antagonist muscle relaxes when the opposite muscle contracts. This has recently been demonstrated in an elegant experiment by Sherrington on the monkey. After section of the third and fourth nerves of the left side, leaving, therefore, the sixth nerve and external rectus muscle alone efficient upon the left eyeball, he finds that appropriate excitation of the cortex gives conjugate movements (*vide* p. 533) of *both* eyes to the right, *i.e.* the left external rectus must have relaxed in consequence of the cortical excitation.

**Movements of the iris.**—Allusion has already been made to the fact that the iris plays the part of a circular muscular diaphragm controlled by nerves. Experimentally, it is found that the central aperture or pupil is diminished by excitation of the third nerve, and by section of the cervical sympathetic—increased by excitation of the sympathetic, and by section of the third nerve.<sup>1</sup> It is usually assumed that the iris is composed of circular or constrictor fibres, and of radiating or dilatator fibres, the third nerve being the motor nerve of the circular muscle, the sympathetic the motor nerve of the radiating fibres. But, owing to the difficulty of demonstrating the anatomical existence of radial fibres, and seeing that direct excitation of an excised iris may cause either elongation or shortening (Grünhagen), room has been left for the introduction of a new hypothesis, to the effect that the iris possesses only a circular or sphincter system of fibres, to which the third nerve is motor and the sympathetic inhibitory. The older view—sympathetic nerve motor to radial fibres—

<sup>1</sup> The eyeball as a whole is rendered less or more prominent by sympathetic section and excitation respectively.



seems, however, to be finally substantiated by the experiments of Langley and Anderson, and of Heese, who observed, and succeeded in recording, radial contraction of the iris by excitation of the sympathetic, after having excised the sphincter, and with a sector of the iris isolated by two radial incisions.

The muscular fibre of the iris is of the non-striated kind (except in birds). Its nerves reach it by way of the ciliary branches of the ophthalmic division of the fifth, which contain fibres derived from that nerve (sensory), from the third nerve (constrictor), and from the sympathetic (dilator). These last have been experimentally traced back through the Gasserian ganglion to the cervical sympathetic, and through the last cervical and first thoracic ganglia to the spinal cord, through the anterior nerve-roots from the fifth cervical to the fifth thoracic—most abundantly through the second thoracic. The region of the cord from which they spring has been termed the *cilio-spinal region* (Budge and Waller), and is subordinate to a bulbar centre above. A few fibres springing from the bulb pass directly to the eye by the channel of the fifth nerve.

Movements of the iris take place in association with accommodation, in response to various stimuli, and in consequence of the administration of certain drugs. Stimulation of the retina by light, or experimental excitation of the optic nerve, causes reflex contraction of the pupil, and on the higher mammalia (dog, cat, and man), with incomplete decussation of the optic nerves, both pupils contract when the retina or optic nerve of one side is stimulated. If one pupil is made to dilate by excitation of the sympathetic, the pupil of the opposite eye contracts, owing to the increased illumination of the first eye. In cases of locomotor ataxy and of general paralysis reflex contraction of the iris in response to illumination, may fail to occur, although the pupil contracts during accommodation (paradoxical reaction, or Argyll-Robertson pupil). On the other hand, in cases of complete blindness due to disease of the cerebral cortex, although there is no *sensation* of light, illumination of the retina may cause reflex contraction of the pupil—*via* optic nerve, basal ganglion (anterior quadrigeminal body?), and third nerve.

Stimulation of the skin, painful impressions, many forms of emotion—such as surprise or awakened attention—muscular exertion, dyspnoea, give rise to dilatation of the pupil; during sleep, on the contrary, the pupil is contracted. After section of the sympathetic, cutaneous stimuli no longer cause reflex dilata-

tion of the now contracted pupil, and the pupil dilates much more slowly on removal of light, but it still contracts with accommodation; after section of the third nerve the pupil is immobile as well as dilated. If the third nerve and the sympathetic be simultaneously and equally excited, the action of the latter

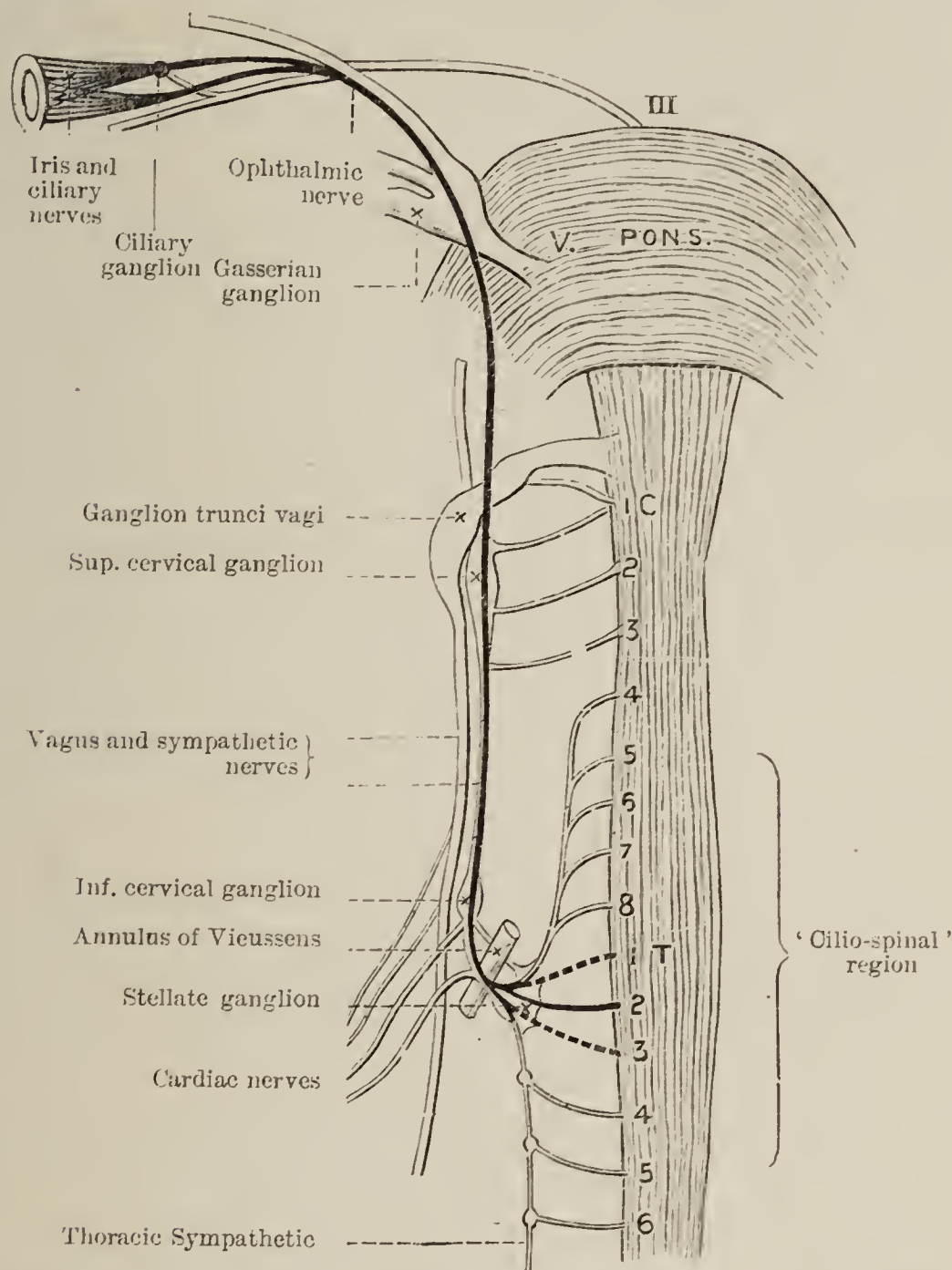


FIG. 250.—DIAGRAM TO ILLUSTRATE THE PATHS OF INNERVATION OF THE IRIS.

Constrictors from the corpora quadrigemina by the 3rd nerve, ciliary ganglion and nerves to the circular muscle of the iris. Dilatators from bulb and cord by anterior roots of the three 1st thoracic nerves, especially the 2nd, rami communicantes, cervical sympathetic and ganglia, Gasserian ganglion, ophthalmic branch of the 5th nerve, ciliary ganglion, and nerves to radiating muscle of iris.

predominates, and the pupil dilates. When death takes place the pupil is for a short time dilated, and subsequently contracted; if the cervical sympathetic is divided on one side before death, dilatation of the pupil on that side does not occur post-mortem. Among drugs, the chief constrictors are morphine and eserine, the chief dilatators are atropine and cocaine. Nicotine



causes a brief dilatation, followed by a prolonged constriction; excitation of the cervical sympathetic now produces no effect, whereas excitation above the superior cervical ganglion dilates the pupil as usual (Langley and Dickinson). Langendorff has shown that at death excitation below the ganglion ceases to be effectual before excitation above the ganglion—*i.e.* nearer to the periphery. Waller showed, in 1853, that the divided cervical sympathetic degenerates up to the superior cervical ganglion, and that four or five days after section its excitation produces no effect on the pupil, while excitation of the ganglion itself produces the usual dilatation. Dilatators of the pupil (*e.g.* atropine) paralyse the ciliary muscle (loss of accommodation); constrictors of the pupil (*e.g.* eserine) cause contraction of the ciliary muscle (spasm of accommodation). The two drugs cause less or more action of one and the same ciliary muscle, and it is very natural to suppose that they cause less or more action of one and the same circular muscle of the pupil; the hypothesis of radial fibres acted upon by atropine is not tenable, for excitation of the third nerve of an atropinised eye produces no effect; the atropine dilatation is due to a paralysed constrictor, and not to a contracted dilatator. This, however, is no proof that a dilatator pupillæ does not exist; the radial contraction of an isolated sector of iris seems to leave no room for doubt.

Movements of the pupil in consequence of variations of light, or of drugs locally applied, will continue to occur in the excised eyes of cold-blooded animals. If a frog's eye be kept for a few minutes alternately in the dark and in the light, the pupil will become obviously larger and smaller.

**Retinal shadows.**—Foreign bodies in the eyeball may, under certain conditions, cast shadows on the retina, and it is possible in various ways to cause the retinal vessels themselves to throw their shadows upon the retina. Such shadows, however produced, excite in consciousness the impression of objects in front of the eye. To determine the presence of foreign bodies focal illumination of the eye should be employed, *i.e.* the light should be placed at that point in front of the eye to which parallel rays emerging from the eye would come to a focus; the eye is thus illuminated by a cylinder of parallel rays from pupil to retina, and a foreign body then causes a retinal shadow, which appears as a dark mass in a bright circular field. By ascertaining the direction and extent to which a given shadow alters its position in the bright field when the light is moved up or down, it is

possible to form a very accurate judgment of the position of the foreign body in the eye. If the foreign body is in the same plane as the iris, its shadow will preserve its relative position in the bright disc. If it is on a plane *anterior* to that of the iris, the shadow will appear to rise and fall as the light is raised or lowered. If it is on a plane *posterior* to that of the iris, the shadow will appear to rise and fall as the light is lowered and raised. In this last case the object appears to move in an opposite sense, while in the second case it appears to move in

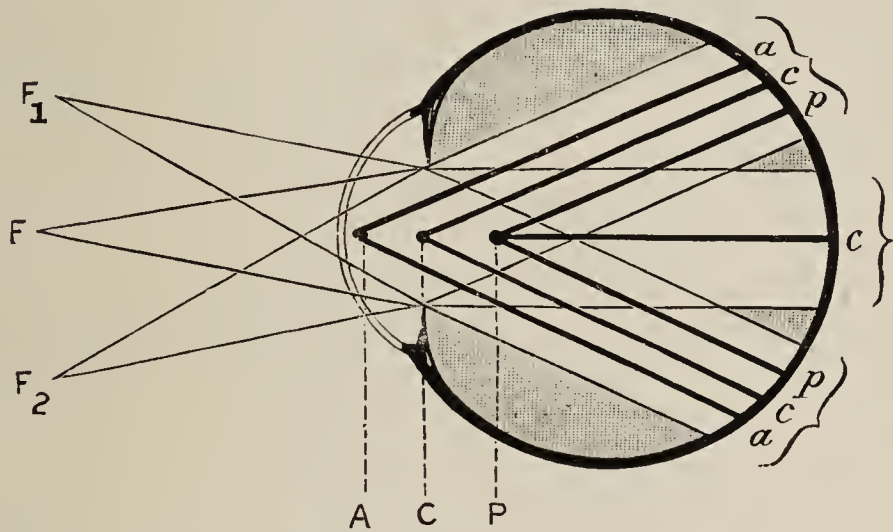


FIG. 251.—POSITION OF FOREIGN BODY ASCERTAINED BY MOVEMENT OF THE SHADOW FORMED WITH FOCAL ILLUMINATION.

—	<i>The foreign body is in front of the plane of the iris, at A.</i>	<i>The foreign body is behind the plane of the iris, at P.</i>
Light raised to $F_1$	Shadow moved towards lower border of retinal disc, <i>i.e.</i> upwards in the bright field.	Shadow moved towards upper border of retinal disc, <i>i.e.</i> downwards in the bright field.
Light lowered to $F_2$	Shadow moved towards upper border of retinal disc, <i>i.e.</i> downwards in the bright field.	Shadow moved towards lower border of retinal disc, <i>i.e.</i> upwards in the bright field.

the same sense, with reference to the direction in which the light is moved (see fig. 251). In any case, the extent of apparent movement is in proportion to the distance of the foreign body from the plane of the pupil; if, for instance, it is very near the retina, a small movement of light may bring the shadow up to or beyond the edge of the bright disc.

Slight shadows caused by small specks or imperfections of the media are common to most eyes; they move with the movements of the eye when attempts are made to look at them; hence their name—*muscæ volitantes*.



Shadows of the retinal vessels may be made apparent in various ways (Purkinje):—(1) A strong light is focussed upon the sclerotic just outside the cornea, the eye being wide open,

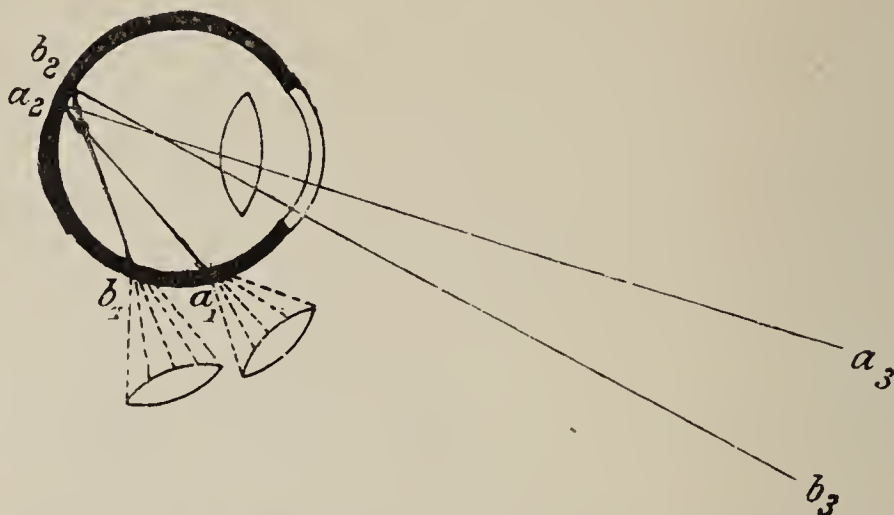
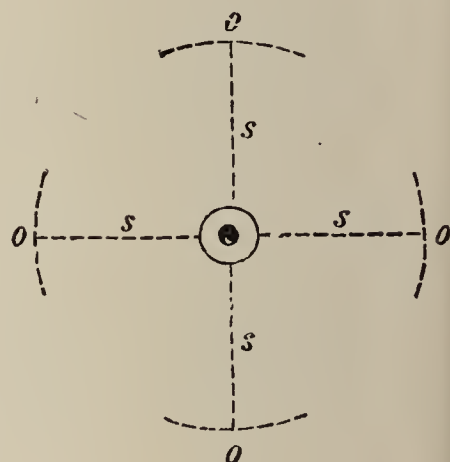
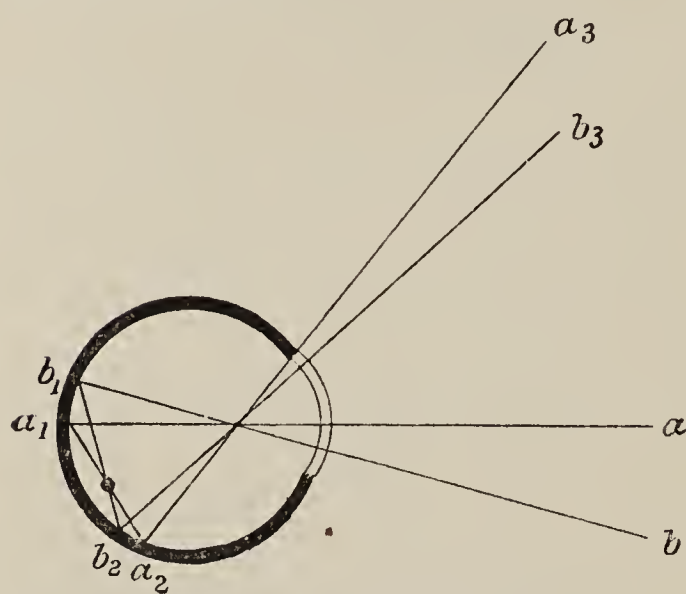


FIG. 252.—PURKINJE'S NETWORK MADE VISIBLE BY ILLUMINATION THROUGH THE SCLEROTIC.

The spot of light,  $a_1$ , gives the shadow,  $a_2$ , of the vessel, which is apparent at  $a_3$ ; if the spot of light is shifted to  $b_1$ , the shadow of the vessel is at  $b_2$ , and is apparent at  $b_3$ , *i.e.* the vessels appear to move in the same direction as the spot of light (Helmholtz).



FIGS. 253 and 254.—PURKINJE'S NETWORK MADE VISIBLE BY ILLUMINATION THROUGH THE CORNEA.

Light from  $a$  is imperfectly focussed at  $a_1$ , where it forms an illuminating spot. A shadow,  $a_2$ , is formed of the vessel, and is referred to  $a_3$  in the field of vision. Considering the diagram to represent a horizontal section of the eye, if the light is moved laterally to  $b$ , the illuminating spot in the eye becomes  $b_1$ , the shadow of the vessel is at  $b_2$ , and the vessel appears to move to  $b_3$  in the field of vision, *i.e.* in the same direction with lateral movement of the light; if, on the other hand, the light be shifted vertically, *e.g.* upwards from  $a$  (above the plane of the paper), the illuminating spot,  $a_1$ , will be lower, the shadow,  $a_2$ , higher, the apparent vessel,  $a_3$ , lower than the plane of the paper, *i.e.* the vessel appears to move against vertical movements of the light (Helmholtz).

More generally stated, so as to include all kinds of positions of light and directions of apparent movement, the relation is as follows: the apparent movement is homonymous when the light is moved in any plane containing the line of vision, and reversed when the light is moved in an arc around the line of vision. If in fig. 254 the circle represents the cornea, its centre the transverse section of the visual line (perpendicular to the paper), a movement of the line along any radius,  $s$   $s$ , will give similar apparent movement; a movement along any portion of the circle  $o$   $o$  will give opposite apparent movement.

directed towards a dark background ; in a few seconds, especially if the light is kept slightly moving, the arborescent pattern of the retinal vessels will be plainly seen. The mode in which a retinal vessel casts its shadow on the retina, and the direction in which the shadow moves and the vessel appears to move, are illustrated in fig. 252.

(2) A readier, though less perfect method of taking cognisance of one's own retinal vessels is to fix the gaze upon a dark background in a darkened room, while a candle is moved slightly up and down or from side to side close to the cheek. The arborescent shadow of the retinal vessels will soon be apparent—most plainly so the horizontal vessels if the candle is moved up and down, the vertical vessels if the candle is moved from side to side. It will be noticeable, moreover, that with vertical movements of the candle held laterally on a level with the eye, the apparent movements of the vessels will be opposite in direction to those of the candle, *i.e.* the vessels appearing to sink in the field as the candle is raised, to rise in the field as the candle is lowered ; with lateral movements of the candle the apparent movements of the vessels will be in the same sense as those of the candle.

The mode in which shadows are formed by this method differs from the preceding one in that the illumination is through the cornea, the light being imperfectly focussed at  $a_1$  (fig. 253), where it forms an illuminating spot analogous to the illuminating spot  $a_1$  in the previous diagram.

(3) If the completely rested eyes are suddenly opened and directed to a white surface, a dark pattern of the vascular network and of the yellow spot will become visible for a few seconds, gradually fading away. This 'entoptic' vision of the yellow spot is due to an absorption of light (more especially of the violet half of the spectrum beyond F) by its yellow spot ; a blue glass or a layer of chrome alum will therefore best make it apparent.

*Phosphenes, or false lights.*—The retina and optic nerve can be excited by mechanical or electrical stimuli ; the sensations to which these give rise are known as *phosphenes*, and are presented subjectively as flashes of light. If the butt-end of a pencil is pressed against the nasal side of the retina while the eye is directed upon obscurity, a circular bright phosphene appears on the temporal side of the field of vision ; similarly, pressure on the temporal, upper or lower sides of the eyeball causes sensations of light referred to the nasal, lower or upper parts of the



field of vision. The sudden relaxation of accommodation may be sufficient to excite the retina mechanically ; the phosphene so produced has the form of a circle. Electrical excitation of the eyeball or of the optic nerve likewise gives rise to phosphenes occupying the whole field of vision, effects which follow the application of either pole of interrupted or of constant currents. Any exact study of polar effects is, however, impracticable, owing to the unavoidable complexity of current-distribution when the electrode is applied to the eye. The flash is, if anything, most pronounced at make when the kathode is on the eyeball, at break when the anode is substituted for it (the second or indifferent electrode being in the mouth). If a fine-pointed kathode is applied to the temporal side of the eyeball, the nasal half of the field of vision of that eye appears bright in comparison with the temporal half ; *vice versa*, if an anode be so applied, the temporal half appears brighter than the nasal half. Even with the eye at complete rest in complete obscurity, excitatory changes are in slight operation ; the field of vision is not dead-black, but faintly illuminated by the so-called ‘intrinsic light of the retina.’

**Objective effects of light upon the retina.**—We have hitherto considered the subjective effects of light acting upon the retina ; there remain to be described certain objective effects of the same cause, which can be appreciated by physical examination. These are :—(1) Bleaching of the ‘visual purple’ ; (2) movements of pigment cells ; (3) movements of cones ; (4) electrical changes.

The effect of light upon the pigment cells of the retina is exactly the opposite to its effect on the pigment cells of the skin. In a retina that has been protected from light before and during removal, the pigment is concentrated in the body of the cells, whereas after exposure to light the pigment is scattered to a considerable distance between the rods—unlike the pigment in the frog’s skin, which is concentrated in the cell under the influence of light, diffused when the skin has been protected from light. The retinal pigment above alluded to has received the names of *melanin* or *fuscine* ; it is the source from which the retinal purple is derived.

In addition to *fuscine*, several other pigments have been identified in the retina, viz. *rhodopsin*, which is confined to the outer part of the rods, and *chromophanes* (red, green, and yellow), which are coloured oil globules situated in the inner segments of the cones. All these pigments are sensitive to light, being bleached by it, but they are not present in all

animals; rhodopsin is not present in the retinae of many birds, chromophanes are not present in mammalia, nor is there any relation between the presence of these pigments and the nocturnal or diurnal habits of various animals.

The pigment of most interest, and that has been most studied, is rhodopsin, or the retinal purple, which can be readily seen colouring the inner surface of the retina of a frog or of a rabbit kept in obscurity for some time previously; the removal of the eye and of the retina should be carried out in darkness, or at most by the aid of a sodium flame. If we expose one eye to ordinary daylight while the other is kept in darkness, the retina of the first eye will be found of a much paler hue, 'bleached,' in comparison with that of the protected eye, the change being much more rapidly brought about in the case of the rabbit's than in that of the frog's eye (Boll, Kühne). If this bleached eye be replaced in darkness, and again examined a short time after, it will be found that the retina has partially or wholly recovered its original pink colour. If, however, this last experiment be repeated with a retina that has been removed from the eye and separated from its layer of pigmented epithelium, the recovery of colour will not take place. From this we learn that the presence of the brown pigment (fuscin) in the outermost layer of the retina is the stock from which is derived the purple pigment (rhodopsin) in the rods.

The retina can be bleached locally in patterns by causing appropriate images of light to be focussed on its surface. By means of a 4 per cent. solution of potash alum, retinal pictures thus produced—*optograms*—can be fixed so as to remain unaltered by the further action of light. The visual purple has been extracted from the retina by means of a  $2\frac{1}{2}$  per cent. solution of bile-salts, as a coloured solution which is sensitive to the action of light (Kühne).

That the retinal purple bears no demonstrable relation to vision is shown by the facts that (1) animals devoid of purple, or moving about in a light so strong that it must have bleached

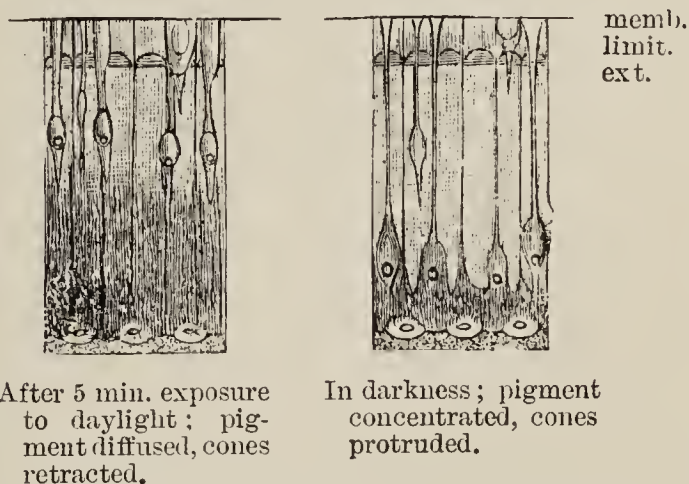


FIG. 255.—ACTION OF LIGHT UPON RETINA.  
(Engelmann.)



every trace of purple, can see ; and (2) retinal purple is absent from the yellow spot, which contains cones only. The popular name 'visual' purple is therefore a misnomer.

Van Stort and Engelmann state that the outer limbs of the cones vary according as preparations are taken from retinae that have been exposed to or protected from light. If the retina has been protected from light, the cones protrude right down to the pigmented layer ; if it has been exposed to light, they are retracted and sessile upon the external limiting membrane. Engelmann states further, that this retraction of the cones is directly controlled by nerves—illumination of one eye producing reflex retraction of the cones of a protected eye, as well as an electrical variation ; strychnia-poisoning, or general faradisation, or dyspnœa, causes the cones of both eyes to be retracted, in the absence of light. The electrical effects of retinal excitation have been described at p. 398.

Dubois has found very interesting structural and functional analogies between the retina and the integument of the siphon of *Pholas dactylus*. An external layer of pigment-cells is connected with a superficial layer of muscular fibres that contract in response to luminous stimuli ; this initial local contraction constitutes a mechanical stimulus of subjacent nerve-cells, and is followed by a secondary total contraction of the longitudinal musculature of the siphon. Histologically, the dermic element excitable by light is composed of (1) pigment-cell, (2) contractile fibre, and (3) nerve-cell. Functionally, the peripheral stimulus is a dermic phosphene, analogous with a tactile phenomenon, viz. a mechanical excitation of the nerve-cell by a movement of the contractile fibre in consequence of a photo-chemical change in the pigment-cell.

**Relations of light perception to retinal stimulation.**—The subjects now to be considered occupy a borderland position between the physiology of the retina and that of the brain. In every subjective study of sensation and in most objective experiments we study in one combined group sense-organ, sensory nerve, and sensory centre. The analysis of any sensation must therefore trench upon cerebral physiology, and that of visual sensation does so in highest degree ; developmentally and anatomically, as well as physiologically, the retina is the most closely cerebral of our sense-organs ; and judgments from visual data form a larger contribution to our daily experience than the judgments derived from all other sensory data put together. We shall in a further chapter make use of these data in general illustration of the

formation of judgments, and shall then give an account of the leading theories into which the facts have been interpreted ; at present we shall simply state these facts as they have been observed, without asserting that they are cerebral, or retinal, or photo-chemical, or psychical. In this allusion to theories we refer particularly to the phenomena of simultaneous contrast, which are described below, and further discussed at p. 565.

The shortest known flash of light, that of the electric spark (of which the duration is less than one millionth of a second), is long enough to produce a sensation, the duration of the latter being very much greater. The sensory effect outlasts its physical cause. Definite numbers expressive of the *duration* of the sensory effect cannot, however, be exactly assigned, as it varies greatly with the strength of stimulation and with the state of the retina.<sup>1</sup> Very strong stimuli give rise to effects so prolonged as to be characterised as after-effects (positive). Many illustrations of this relation between stimulus and sensation are matters of everyday experience. (1) A luminous body, *e.g.* a firebrand, rapidly whirled round, causes the sensation of a fiery circle ; different retinal points are excited in such rapid succession that the first-made sensations are still in force when the last-made sensations appear. (2) A disc of two colours, if rapidly rotated, is seen of a uniform resultant colour ; *e.g.* if the two colours are complementary the resultant is in the direction of white ; the same retinal points are alternately excited by the two colours in such rapid succession that the sensations of both co-exist and are fused in consciousness. (3) A succession of stimuli such as that produced by the rotation of a black disc with white sectors, produces a sensation analogous with muscular tetanus, *i.e.* infrequent stimuli are perceived separately, more frequent stimuli are perceived incompletely fused, and, finally, stimuli above a given limit of frequency give rise to a perfectly uniform sensation ; this last is analogous with complete tetanus, the incompletely fused ‘flickering’ sensation with incomplete tetanus or muscular clonus, the separate sensations with separate muscular contractions. If the white sectors are strongly lighted, the *minimum frequency* may reach 50 stimuli per second before complete uniformity of sensation is reached ; with a weak light, the minimum limit may be as low as 10 per second. Charpentier’s experiments bring out a further analogy between retinal and muscular excitatory

<sup>1</sup> It would be more correct to say ‘of the retino-cerebral apparatus’ ; but the abbreviation is sanctioned by long use.



phenomena. According to his measurements, a local retinal stimulus gives rise to an oscillatory process with a duration of about 0·028 sec., a frequency of about 36 per sec., and, in relation to the retina, with a propagation-velocity of 72 mm. per sec.

*Strength* of stimulus and strength of sensation are not directly proportional. With strong stimulation a larger increment is requisite to excite the sensation of difference than with weak stimulation ; or, to use a concrete instance, the difference of one candle between two lights of 9 and 10 candle-power respectively will be very obvious, while there will be no perceptible difference between two lights of 1,000 and 1,001 candle-power. Otherwise expressed, the amount of stimulus requisite to excite a perception of difference between two stimuli bears a constant relation to their magnitude, which can be expressed in the form of a fraction. The value of this fraction in the case of light is  $\frac{1}{100}$ , *i.e.* we can see a difference between two lights of 10 and 10·1, or of 100 and 101, or 1,000 and 1,010 candle-power. A difference of one candle is an obviously large difference when the lights are of the first-named magnitude, it is a just perceptible difference when they are of the second-named magnitude, it is quite imperceptible between lights of the third-named magnitude. A single brief stimulus, such as an electric spark or a flash of lightning, often appears to be reduplicated, *i.e.* gives rise to a double sensation.

*After-images.*—We have alluded to positive after-images as being simply an instance of after-sensation ; they are called ‘ positive ’ because their lighting is similar to that of the original visual images, and in contradistinction from a second variety of after-images, in which the shades of the original picture are reversed ; images of this kind are negative after-images, which depend for their production upon retino-cerebral fatigue. They may easily be elicited. If, for instance, a well-lighted white pattern on a black ground be steadily gazed at for 10 to 15 seconds, a negative after-image consisting of a dark pattern on a light ground will appear when the gaze is transferred to any uniform surface. Negative after-images are likewise produced by coloured patterns, and in this case the colour of the after-image is complementary to that of the original object ; *e.g.* the after-image of a red pattern is green ; of green, red ; of blue, orange ; of orange, blue ; &c.

The following after-image experiment shows that an apparently completely saturated colour may be subjectively more fully saturated. A small green disc is gazed at for a few seconds, the

eyes are then turned towards a saturated red ground ; the red after-image of the green disc is redder than the surrounding red field. On the Young-Helmholtz theory this is attributed to fatigue of the cones excitable by green, and the consequent greater purity of the red excitation in the retinal area which has been covered by the green image. Similar results have been obtained with pure spectral colours, and lead to the conclusion that the most saturated objective colours do not with the non-fatigued eye produce the maximum sensation of saturated colour.

Negative after-images are the most obvious sign of retinal—or, properly speaking, of retino-cerebral fatigue ; another sign of this process can be appreciated during the contemplation of objects. An object that is just visible in a dim light may be lost sight of if an attempt is made to fix the eyes upon it ; at nightfall the horizon between sea and sky may fade out of sight when a prolonged attempt is made to define it, although the line of separation may remain quite perceptible by indirect vision. A star of just perceptible magnitude may become invisible as an attempt is made to keep it in view, though it reappears as soon as the line of vision is diverted by a few degrees. This last observation, well known to astronomers, is probably to be explained as due to the dull excitability of the fovea centralis of the retina. All the observations above cited show that a stimulus of constant intensity produces, if prolonged, a diminishing excitation.

**Contrast.**—A sheet of ‘white’ note-paper appears very white against a black background, but far short of white against fresh-fallen snow. A piece of red cloth appears redder on green turf, less red against a red background. A yellow skin looks white by the yellow light of gas. These are instances of contrast and of absence of contrast between simultaneous impressions, in consequence of which white, red, and yellow appear to be more or less white, red, and yellow according to circumstances.

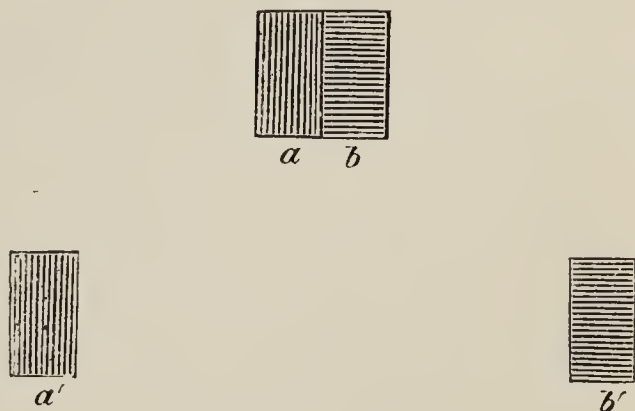


FIG. 256.

A simple but rough way of studying contrast is that which was first adopted by Chevreul. Two slips, *a* and *b*, of coloured paper are fixed side by side ; two similar strips, *a'* and *b'*, are fixed at a



distance from the first pair. Comparing  $a$  with  $a'$ , we see an apparent difference, which is due to the influence on the sensation from  $a$  of the sensation from the adjacent slip,  $b$ . Similarly, by comparing  $b$  with  $b'$  we are made sensible of the influence of  $a$  upon  $b$ . Each colour,  $a$  or  $b$ , will appear as if mixed with a certain proportion of the complement of the other colour,  $b$  or  $a$ , *i.e.* their difference is increased.

The experiment of coloured shadows further illustrates this principle. A rod is fixed so as to throw upon a white screen the shadows (1) of daylight admitted through a hole in a shutter, (2) of candlelight. The first light is white and illuminates the screen with the exception of the portion occupied by the first shadow ;

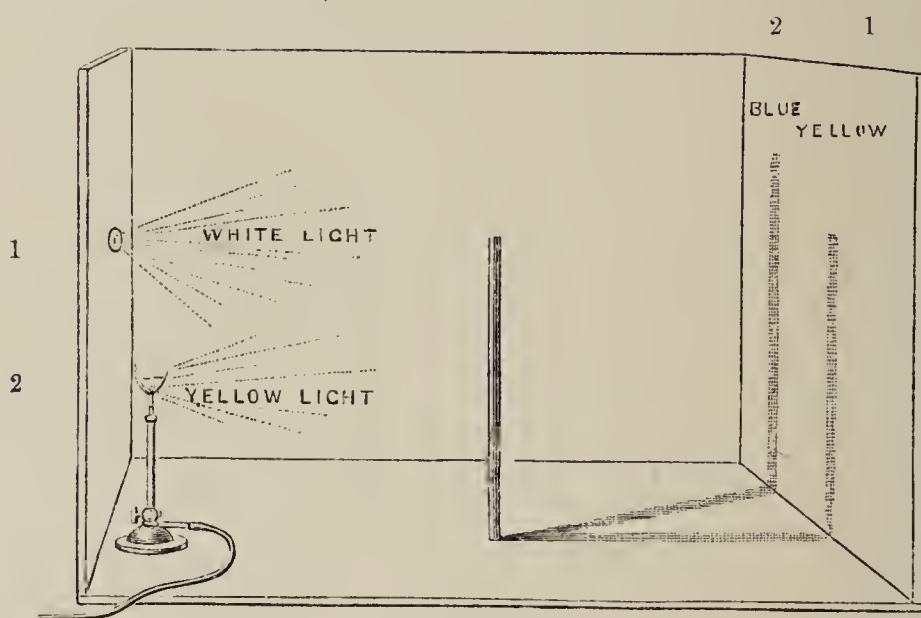


FIG. 257.

the second light is yellow and illuminates the screen with the exception of the second shadow ; so that light 1 (white) illuminates shadow 2, light 2 (yellow) illuminates shadow 1. Objectively, then, shadow 2 should be grey, shadow 1 should be yellow, and the rest of the screen should be pale yellow.

As seen, however, shadow 2 is not grey, but *blue*, in contrast with the yellow screen and still more yellow shadow 1.

Or again : Four candles placed in a row illuminate a white screen in a dark room. A second screen is partly interposed between the candles and the first screen, on which we therefore have four vertical strips, illuminated by 1, 2, 3, and 4 candles respectively. Each strip is uniformly illuminated throughout, yet it does not appear so. The border of a strip nearest to a darker neighbour looks lighter than the border nearest to a lighter neighbour.

Contrast colours are far more effectively excited by pale than by saturated colours. This is easily verified by the following

experiment (H. Meyer). A patch of grey paper placed on a sheet of green paper appears little or not at all altered in tint, but if both papers are covered with thin white tissue-paper, whereby the green colour is much weakened, the grey patch at once assumes the contrast tint, pale red. This result, which is at first sight very surprising, illustrates the fact that differences of sensation are much greater with weak than with strong stimulation, so that any error in the sensory estimate is exaggerated when the sensation and the contrasted counter-sensation are weak. Saturated colours in a painting modify greys and whites less than do pale colours.

All the examples given above are instances of *simultaneous* contrast, in which the exciting cause in one portion of the retina modifies cerebral sensations derived from a different portion of the retina, the compared lights or colours being viewed simultaneously. But under ordinary circumstances the eye wanders, so that the effect of simultaneous contrast is reinforced by negative after-effects, which constitute the phenomenon of *successive* contrast. In this case colours fall in succession upon the same portion of the retina ; each colour modifies the subsequent colour in this sense, that the second colour appears as if mixed with a certain proportion of the complement of the first. The consequence of the first excitation is a negative after-effect or after-image, which becomes fused with the second excitation. As already stated, the after-effect of green is its complement, red, so that if, after looking at a green surface, we look at any other colour, that colour appears to contain less green or more red. The effect in successive contrast is thus simply what we are already familiar with under the name of negative after-image.

We are very liable to get effects in which successive and simultaneous contrasts both take part ; it is extremely difficult to keep the eye steady for many seconds at a time ; the eye wanders, and we then obtain the contrasting excitations successively in the same part, instead of simultaneously on different parts of the retina. Finally, the observation should be made that a negative after-image, as a successive contrast effect, may give rise to a surrounding effect negative to itself as a simultaneous contrast effect. A green pattern gives a red after-image on a greenish background ; the red after-image being the complement of the green pattern (successive contrast), the greenish background being the complement of the red after-image (simultaneous contrast).



Thus we have recognised three varieties of contrast :—(1) Simultaneous contrast with the eye fixed (*a*) between two retino-cerebral effects, or (*b*) between two retino-cerebral after-effects ; (2) successive contrast, with the eye moved, as a retino-cerebral effect superposed upon a retino-cerebral after-effect.

*Binocular effects.*—It is possible to obtain a contrast effect upon one eye by the chromatic stimulation of the other eye. A slip of white paper on a black background and seen double, with a blue glass in front of the right eye and a grey glass in front of the left eye, appears blue to the former and yellow to the latter. It is also possible, with the adoption of suitable precautions, to obtain the mixture of two colours separately presented to each eye. To this end it is necessary that the coloured surfaces should be perfectly even and unlimited by any outlines. The mixture is less perfect than that which is obtained by causing the two colours to coincide upon the same retina, and at the first attempts to observe it we are liable to alternate attention to one or the other eye, and see almost exclusively one colour or the other, or a patchwork giving the effect of a polished surface ; but even at its best the binocular mixture is less complete than the monocular effect, and always darker (Fechner, Hering).

*Observations with slowly-revolving discs.*—In a strong light (direct sunlight) several very beautiful experiments may be made with the aid of slowly-revolving discs ; of the following four, the first is simply a contrast experiment in a striking form, the next three are demonstrative of oscillatory processes taking place in the retina ; these are of a physiological nature, and much slower than the physical oscillations of light.

1. A disc,  $\frac{3}{4}$  coloured and  $\frac{1}{4}$  white, as in fig. 216, revolving slowly, once in one or two seconds, gives on the white quadrant the complementary colour if the centre of the disc is steadily regarded. This, of course, is a successive contrast effect with a moving surface in front of the fixed eye, and the contrast colour comes out better in this way than in any other.

2. A similar disc,  $\frac{3}{4}$  black and  $\frac{1}{4}$  white, similarly regarded, gives on the white quadrant a black band or sector, not far from the black edge that has just passed in front of the eye. This is a rebound effect indicating an oscillatory process in the retino-cerebral apparatus ; the first effect at the arrival of the white border is white, but this first effect is followed by an after-effect that is black ; estimating from the speed of revolution and the apparent position of the black band in the white, this black after-effect is found to follow the main black effect at an interval of about .015 sec. (Charpentier), *i.e.* the duration of the

retinal oscillation is about  $\cdot 03$ , and the frequency thus indicated would be 33 per second. It is owing to such an oscillation that a single bright stimulus gives a double bright sensation.

3. A black disc with narrow open spaces, revolving in front of the eyes fixed upon a white surface lighted by the sun, gives rise to a magnificent lilac field when the rate is such that the retina is stimulated between 33 and 66 times per second (practically between 40 and 60), *i.e.* when each stimulus occurs during the negative after-effect of the preceding stimulus. At higher and at lower frequencies the field is colourless, and at the optimum frequency a yellowish centre is seen to the lilac field, corresponding to the spot of distinct vision, where the retinal purple is deficient. Charpentier, to whom we owe this observation, considers the coloured sensation to be due to entoptic vision of the retinal purple.

4. A large black disc with a small white patch near its circumference, revolving, say, twice per second, gives to the observer a sensation of a white *ribbed* streak occupying a portion of the black surface. This ribbed appearance is a sign of an oscillatory process in the retino-cerebral apparatus; the interval between two ribs diminishes as the rate of revolution increases, and measurements made at different rates indicate that the excitatory state propagates itself from its point of origin with a retinal velocity of about 72 mm. per sec., and a wavelength of 2 mm.

**Animal luminosity.**—Many animals (and plants), *e.g.* fireflies and glow-worms, are phosphorescent, *i.e.* they produce luminous matter. The physiology of this phenomenon has been very fully analysed by Dubois, who experimented upon insects (*Elateridæ*) and upon a mollusk (*Pholas dactylus*). He finds that the phosphorescent material is a granular protoplasm-product elaborated by cutaneous glandular organs under the control of nervous ganglia. In analogy with the action of diastase upon dextrin as regards sugar-production, he refers the production of the final luminous matter to a ferment action between two substances—a ferment, 'luciferase,' and a crystalline body, 'luciferin'—which he has separated from *Pholas dactylus*. He has experimentally reproduced a luminous material by mixing the two non-luminous solutions of these bodies; spreading one solution upon paper, he traces upon it with the second solution a pattern that becomes luminous; or again, placing the mollusk in sulphuric ether, he obtains an extract that separates into two layers—an upper, ethereal layer, containing the luciferin, and a lower, aqueous layer, containing the luciferase, both layers being non-luminous, but when shaken giving a luminous band at their surface of contact. He gives, *e.g.* the photometric value of the light from one glow-worm equal to about  $\frac{1}{40}$  candle. Experimenting on *Elater noctilucus*, Fuchs was able to make its luminous organs light up by electrical excitation of their nerves.



## CHAPTER XIII

## SOUND AND HEARING

PAGE

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Noise—Musical sound—Strength—Pitch—Timbre—Speed—Wave-length—Resonation—Consonance of strings or membranes—Overtones or upper partial tones—Beats—Undertones or beat-tones—Concord and discord.

## 467 The Ear.

External, middle, and internal ear—Membrana tympani—Ossicles—Tensor tympani—Stapedius—Eustachian tube—Hearing *via* the cranium.

## 470 The internal ear—Semicircular canals—Cochlea—Membrana basilaris—Organ of Corti—Hair-cells—Theories.

## OTHER SENSES

## 475 Taste—Smell—Touch—Cutaneous sensation.

## SOUND

**Physical data.**—*Sound* is a sensation produced in the organ of hearing by vibrating bodies. Vibrations may be seen, felt, and *heard*; when they are heard, we call their resultant a *sound*. Irregular vibrations cause noises, regular vibrations cause musical sounds.

Musical sounds or tones differ in *intensity* or *strength*, in *pitch*, and in *timbre* or *character*. Intensity depends upon amplitude of vibration, pitch depends upon the number of vibrations per unit of time, timbre depends upon upper partial tones, which accompany a fundamental tone.

Sounds of all strengths travel at the rate of 1,100 feet (330 meters) per second in air of 15° Centigrade. Given the speed at which sound travels and the number of vibrations per second, it is easy to calculate the *wave-length* of a single vibration. Thus, the wave-length in a note of 11 vibrations per second will be 100 feet; in a note of 55 vibrations, it will be 20 feet; in a note of 100 vibrations, it will be 11 feet.

A note *resonates* in a cylinder having the same length as the wave-length of vibrations that constitute the note, or having a length which is an exact divisor of the wave-length of the original note. If

different notes are simultaneously sounded, a cylinder will resonate with and reinforce the note of corresponding wave-length, and that note will be heard louder than the accompanying notes. Similarly, tense strings and membranes will *resonate* with certain notes rather than with others, according as the vibration-numbers of the notes do or do not correspond with the vibration-number of the string or membrane when pulled or struck. A given note loudly sounded in front of a series of strings tuned to different notes, will set in vibration certain strings, to the exclusion of others.

*Overtones, or upper partial tones.*—The principal or fundamental vibrations constituting a tone are usually accompanied by secondary tones, higher and of less intensity, which greatly affect its ‘timbre.’ A simple tone free from overtones is soft, weak, and dull. The tone of an instrument giving overtones in the chord of that tone is fuller, richer, and more sonorous than a simple tone. If the overtones are uneven, but not excessive, the musical character of the tone remains, and may be peculiarly agreeable; but if the uneven overtones are too marked, or too numerous, the tone is rendered rough or penetrating, and is no longer agreeable to a civilised ear.

*Beats; undertones, differential, or beat-tones.*—Two slightly different notes, sounding together, alternately strengthen and weaken each other by interference, and the succession of phases is audible as *beats*. A tuning-fork of 436 vibrations sounding with a tuning-fork of 440 gives 4 beats per second, *i.e.* the frequency of beats per second corresponds with the difference in the numbers. Beats, in sufficiently rapid succession, give rise to a secondary, differential, or beat-tone. Thus, a tone of 132 vibrations with a tone of 198 vibrations gives a beat-tone of 66 per second. Widely different notes can also give beat-tones, the vibration-frequencies of which will correspond with the ‘positive’ and ‘negative’ remainders obtained by dividing the vibration-frequency of one note by that of the other (König). Notes with vibration-frequencies of 110 and 440, sounding together, give no beat-tone, there being no remainder on dividing 440 by 110. Notes of 120 and 440 will give beat-tones with vibration-frequencies of 80 and 40.<sup>1</sup>

Tones are *concordant* when their vibration-numbers bear to each other simple ratios, expressible by small whole numbers. Such tones harmonise and form concords or chords when sounding together. Tones are *discordant* when their vibration-numbers do not bear to each other ratios as above. Such tones clash and form discords when sounding together.


In the major chord	. .	C	E	G	c
the vibration-numbers are		132	165	198	264
their ratios are	. . . .	4	5	6	8

<sup>1</sup> 120 goes into 440 three times, leaving 80 as the *positive* remainder; or four times all but 40, which is the *negative* remainder.



In the discord . . . . C D E  
the vibration numbers are 132 148·5 165  
and are not reducible to small whole numbers.

In forming numerical estimates of auditory sensibility, it is necessary to remember:—(1) that the strength of sound made by a body allowed to drop upon a surface is proportional to the square root of the height of drop; (2) that the strength of sound varies inversely as the distance squared between origin and ear; *e.g.* the strengths of sound made by a ball falling from heights 1, 4, 9, 16 are 1, 2, 3, 4; the strengths of a given sound at distances 1, 2, 3, 4 are 1,  $\frac{1}{4}$ ,  $\frac{1}{9}$ ,  $\frac{1}{16}$ .

						
B <sub>,,</sub> 61 $\frac{7}{8}$ (60)	B <sub>,</sub> 123 $\frac{3}{4}$ (120)	B 247 $\frac{1}{2}$ (240)	b 495 (480)	b <sub>,</sub> 990 (960)	b <sub>,,</sub> 1980 (1920)	b <sub>,,,</sub> 3960 (3840)
A <sub>,,</sub> 55 (53·3)	A <sub>,</sub> 110 (106·6)	A 220 (213·3)	a 440 (426·6)	a <sub>,</sub> 880 (853·3)	a <sub>,,</sub> 1760 (1706·6)	a <sub>,,,</sub> 3520 (3413·3)
G <sub>,,</sub> 49 $\frac{1}{2}$ (48)	G <sub>,</sub> 99 (96)	G 198 (192)	g 396 (384)	g <sub>,</sub> 792 (768)	g <sub>,,</sub> 1584 (1536)	g <sub>,,,</sub> 3168 (3072)
F <sub>,,</sub> 44 (42)	F <sub>,</sub> 88 (85·3)	F 176 (170·6)	f 352 (341·3)	f <sub>,</sub> 704 (682·6)	f <sub>,,</sub> 1408 (1365·3)	f <sub>,,,</sub> 2816 (2730·6)
E <sub>,,</sub> 41 $\frac{1}{4}$ (40)	E <sub>,</sub> 82 $\frac{1}{2}$ (80)	E 165 (160)	e 330 (320)	e <sub>,</sub> 660 (640)	e <sub>,,</sub> 1320 (1280)	e <sub>,,,</sub> 2640 (2560)
D <sub>,,</sub> 37 $\frac{1}{8}$ (36)	D <sub>,</sub> 74 $\frac{1}{4}$ (72)	D 148 $\frac{1}{2}$ (144)	d 297 (288)	d <sub>,</sub> 594 (576)	d <sub>,,</sub> 1188 (1152)	d <sub>,,,</sub> 2376 (2304)
C <sub>,,</sub> 33 (32)	C <sub>,</sub> 66 (64)	C 132 (128) (8-foot organ pipe)	c 264 (256) (middle c)	c <sub>,</sub> 528 (512) (standard tuning- fork)	c <sub>,,</sub> 1056 (1024)	c <sub>,,,</sub> 2112 (2048)

Musical notes and their vibration-frequencies. The numbers in parentheses are such as would be obtained by taking the series C<sub>,,</sub> . . . . c<sub>,,,</sub> in successive powers of 2, viz. . . . 2<sup>n</sup>.

## HEARING

The organ of hearing is composed of: (I.) a receptive part—the external, middle, and internal ear; (II.) a transmissive part—the auditory nerve; (III.) a perceptive part—the cortex of the brain.

**The ear.**—Anatomically, the ear consists of: (1) the external ear, comprising the pinna and auditory meatus, by which, in normal hearing, sonorous vibrations are conducted to and concen-

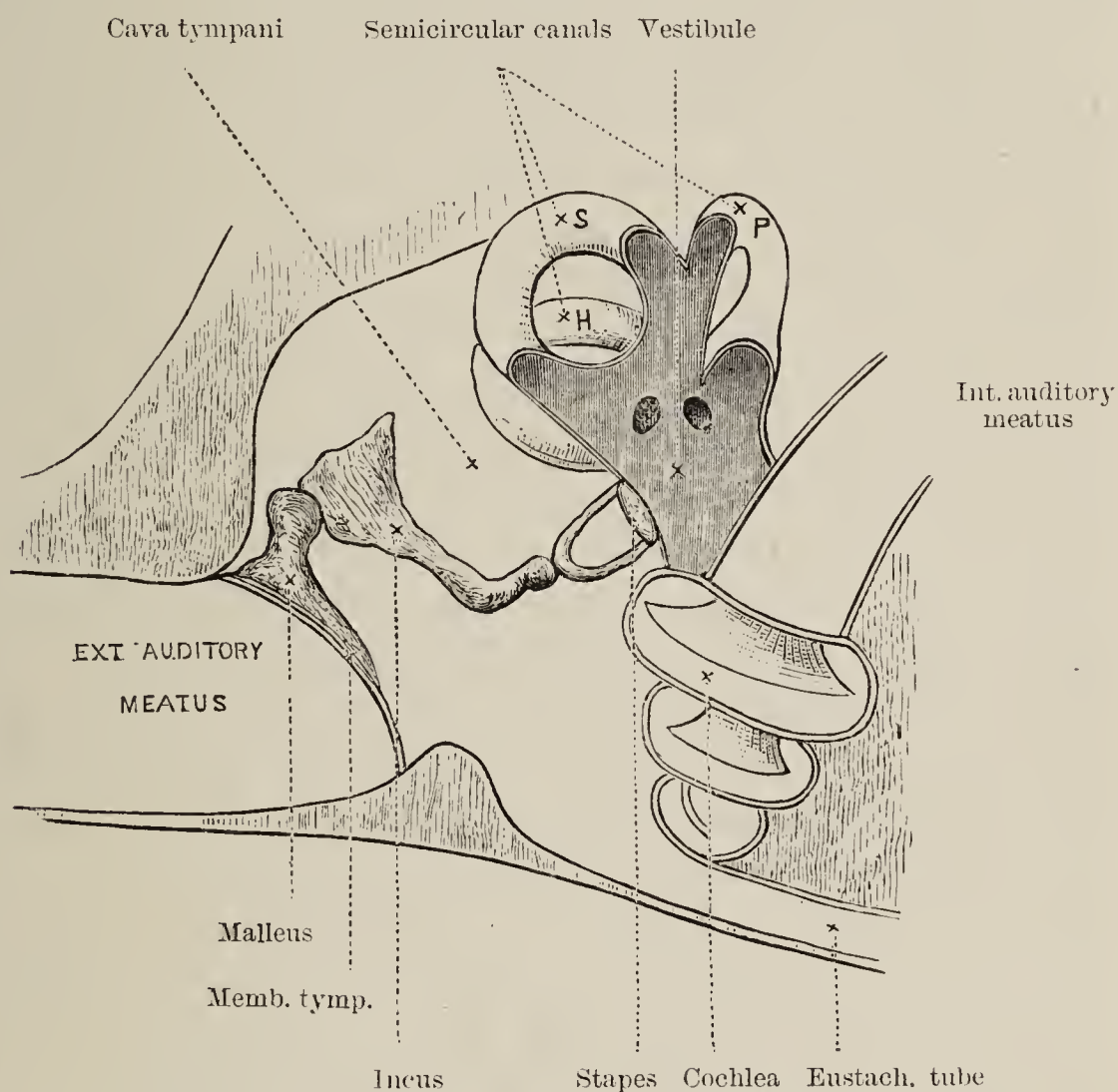


FIG 258.—DIAGRAMMATIC VERTICAL SECTION THROUGH THE AUDITORY APPARATUS.

trated upon the tympanic membrane or drum, which divides the external from the middle ear. (2) The middle ear, or tympanum, containing a chain of small bones, the three auditory ossicles, malleus, incus, and stapes. (3) The internal ear, comprising the vestibule, cochlea, and semicircular canals, and containing the peripheral termination of the auditory nerve—the hair-cells of the organ of Corti.

Sonorous vibrations arriving at the membrana tympani cause that membrane to vibrate; its vibrations are transmitted through the auditory ossicles to the fluid filling the internal ear, and the vibrations of that fluid excite the peculiarly modified



epithelium, from which the auditory nerve is the conducting channel to the brain.

The *membrana tympani* possesses a physical peculiarity distinguishing it from ordinary stretched membranes, which resonate with particular notes, inasmuch as it does not exhibit this physical preference to any marked degree, but vibrates with equal readiness to all kinds and tones of sound. It may, nevertheless, in some degree be attuned or accommodated to differences of pitch by means of muscles, which pull upon the auditory ossicles, and through the Eustachian tube by variations of air-pressure.

*The ossicles.*—The vibrations of the membrane are transmitted along the ossicles, not merely as vibrations of sound are transmitted along solids, but as actual movements of the bones, acted upon and acting as levers. The movements are not molecular, but molar, and they have been observed and measured. The handle of the malleus attached to the *membrana tympani* forms one end of the lever—that to which the force is applied; the base of the stapes, attached by its orbicular ligament to the margin of the fenestra ovalis, forms the other end of the lever—that at which the force effects movement; the hinge or axis of rotation is in an antero-posterior line joining the points of attachment of the malleus and incus to the wall of the tympanic cavity. The relation between the two arms of this lever is such that the oscillation of the tympanic membrane is transmitted to the base of the stapes with an amplitude diminished to two-thirds, and with corresponding mechanical advantage. The amplitude of movement of the tympanic membrane ranges from a maximum of  $\frac{1}{10}$  mm. down to the immeasurably small; that of the stapes, from a maximum of  $\frac{1}{15}$  mm., likewise down to the immeasurably small—or, to give some idea of how small, down to less than  $\frac{1}{100000} \mu$ .

The *tensor tympani* muscle, attached to the handle of the malleus, pulls the latter, and with it the tympanic membrane, directly inwards, thus tightening the membrane. It is served by the fifth nerve. With this movement the stapes is more closely applied to the fenestra ovalis and tilted forwards; amplitude of vibration is thus limited, and sounds, especially low notes, are consequently weakened. The *stapedius* muscle, attached to the neck of the stapes, tilts the latter backwards. It is served by the seventh nerve. Its influence upon the transmission of sound is not known.

The *Eustachian tube* establishes communication between the tympanic cavity and the pharynx, and thus admits atmospheric

pressure to the cavity of the tympanum. It is probably not continuously open, but only so on more or less frequent occasions, *e.g.* during the act of swallowing. If it be obstructed, atmospheric pressure on the outside of the membrana tympani is not balanced by counter-pressure from the inside, the tympanic cavity becomes filled by exudation, and hearing is interfered with. Hardness of hearing from this cause is temporarily removed by inflation of the tympanum; this may be effected by a strong expiratory effort of the patient with mouth and nostrils closed; or, the patient being directed to swallow, air may be forced through from the nostrils by an operator during the act of deglutition. Hearing is temporarily dulled whenever air-pressure within the tympanic cavity is less or greater than that upon the outer surface of the membrane; during a forcible inspiration or expiration with closed mouth and nostrils, sounds, especially low notes, are less distinctly heard than when tympanic air-pressure is normal, *i.e.* equal to atmospheric pressure.

*Hearing via the cranium.*—If, while the vibrations of a tuning-fork are dying away and have just become inaudible by the usual channel, the handle of the fork be applied to the teeth or skull, the sound will become again distinctly audible—*i.e.* the vibrations are transmitted by the skull. If, while the tone is being listened to in this manner, one ear be gently closed, the sound will be intensified on that side. A fully satisfactory explanation of this fact cannot be given; it was formerly supposed that the vibrations were transmitted by the bones of the skull directly to the internal ear; a sound *via* the cranium more distinctly heard on one side (that side being defective to sounds reaching it by the usual channel) than on the other, was believed to indicate defect of the sound-conducting apparatus (tympanic membrane and ossicles). Both these suppositions have been disproved—the vibrations heard through the skull are transmitted by its bones to the membrana tympani and ossicles, and thus onwards to the internal ear, but they have not been proved to be directly transmitted to the internal ear better than through the middle ear. Defects of the middle ear and of the internal ear alike interfere with the hearing of sounds *via* the cranium, as well as *via* the meatus; if a person who is dull of hearing hears a tuning-fork on the cranium louder in an affected ear, he probably has a blocked meatus, and not a defect of the middle ear; if he had the latter, the tuning-fork would be less audible on that side.

The apparatus of the middle ear—membrana tympani and



auditory ossicles—is useful, but not absolutely indispensable ; hearing is greatly impaired by their destruction, but it is not by any means abolished, and slight perforations of the membrana are common without obvious hardness of hearing. Some small proportion of sound must therefore be transmissible through the cranium to the terminal apparatus of the internal ear. It has even been stated that hearing is possible after destruction of the internal as well as of the middle ear ; Ewald found that pigeons, after extirpation of the labyrinth on both sides, reacted to sounds—*i.e.* seemed to hear—and he was of opinion that in such case vibrations through the cranial bones directly excited the auditory nerve.

**Physiological anatomy of the internal ear.**—The internal ear, or labyrinth, consists of the vestibule, the cochlea, and the three semicircular canals. Of all these parts the walls are bone, the lining is membrane, the contents are fluid (endolymph and perilymph), and a modified epithelium, which constitutes the commencement of the auditory nerve ; otoliths and so-called ‘sand,’ composed of calcium carbonate, also form part of the contents of the labyrinth, but have no clearly assignable function. Of the two principal branches of the auditory nerve—*cochlear* and *vestibular*—the latter is connected with the function of equilibration, the former is the nerve of hearing ; entering the base of the cochlea, it spreads out as a spiral sheet of fibres interrupted by a spiral ganglion ; above the ganglion these fibres are medullated, beyond it they are non-medullated, finally terminating in a fine network of fibrils round the hair-cells of the organ of Corti. The cochlear nerve-fibres are of smaller calibre than the vestibular nerve-fibres ; both kinds are medullated.

*Functions of the internal ear.*—As regards the *vestibule*, there are no facts to justify any dogmatic statement whatever ; at the most we may regard as not impossible that its epithelium may be excitable by vibrations of the endolymph. As regards the *semicircular canals*, we have reason to believe that they have to do with equilibration, and have quite dismissed the gratuitous supposition that they are concerned in the appreciation of the direction of sound. The cochlea is evidently the most highly specialised portion of the internal ear ; its minute anatomy and its analogies with well-understood physical instruments, indicate with reasonable certainty that it constitutes the physiological receiver of sonorous vibrations—and there are a few physiological and pathological facts in additional support of this conclusion.

The cochlea contains the terminal apparatus of the auditory nerve. It consists essentially in a spiral tube of bone, 25 to 30 mm. long, lined by membrane, and divided by membranous partitions into three channels—*scala vestibuli*, *scala tympani*, and *canalis cochleæ*; the lamina spiralis ossea and the *membrana basilaris* divide the *scala tympani* from the *scala vestibuli*, and this membrane forms the floor of the *canalis cochleæ*, of which the roof is formed by the membrane of Reissner; the *scala vestibuli* commences at the fenestra ovalis, and communicates with the *scala tympani* at the helicotrema; the *scala tympani* commences at the fenestra rotunda; the fenestra rotunda is covered by membrane; the fenestra ovalis is covered by the base of the stapes and its orbicular ligament, so that any vibration

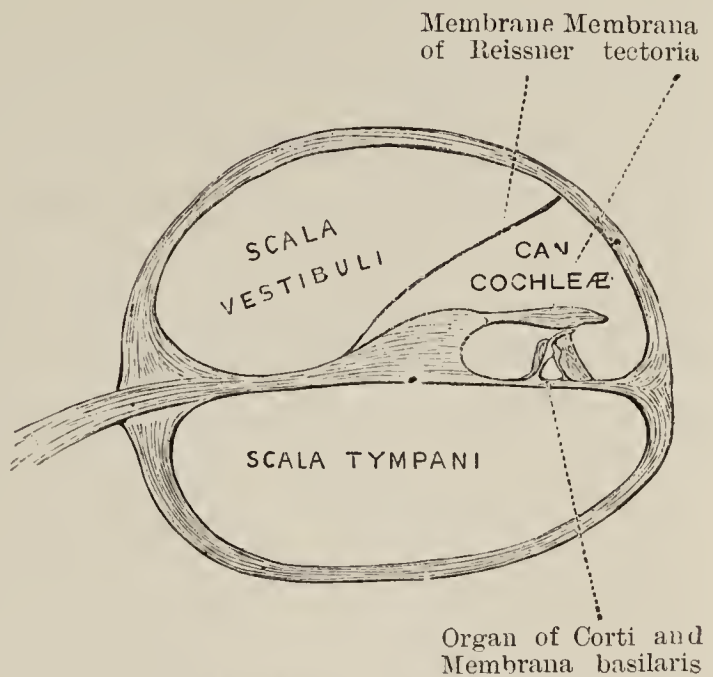


FIG. 259.—DIAGRAMMATIC TRANSVERSE SECTION THROUGH A TURN OF THE COCHLEA.

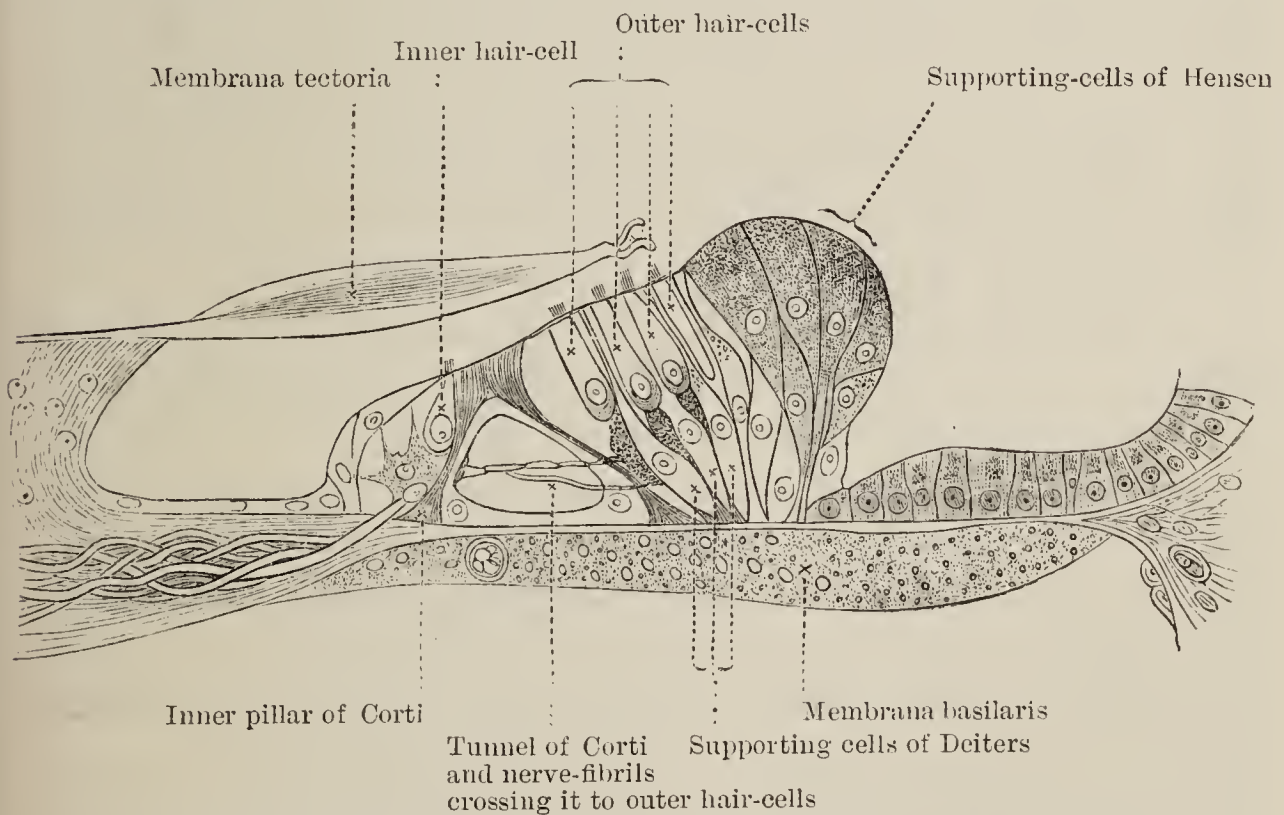


FIG. 260.—TRANSVERSE SECTION THROUGH THE ORGAN OF CORTI; MAN. (Retzius.)

from the *membrana tympani*, transmitted along the ossicles, must pass through the entire length of the cochlea. The floor of the *canalis cochleæ*, or *membrana basilaris*, carries



an avenue of cells which constitute the *organ of Corti*; this consists of a double row of stiff cells, the inner and outer rods of Corti, and of several rows of *hair-cells*—a single rank of inner hair-cells, and a border of outer hair-cells three or four deep, to which the terminal filaments of nerve-fibres have been traced. The organ of Corti is covered by a thin reticulated or fenestrated membrane, through which project the hairs of the hair-cells, and by a loose thick membrane, the *membrana tectoria*, that rests upon the organ. According to Helmholtz, it is of importance to recognise that the *membrana basilaris* is equivalent to a series of radial fibres

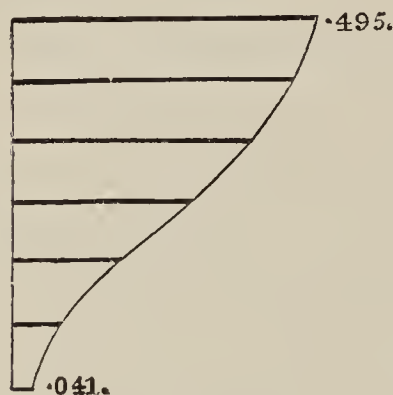


FIG. 261.

Diagram to show the manner in which the breadth of the *membrana basilaris* increases from below upwards. (After Hensen's measurements.)

stretching from the margin of the spiral lamina to the spiral ligament, and that the breadth of the membrane, *i.e.* the length of the fibres, increases from below upwards—at the base of the cochlea  $41\ \mu$ , at its apex,  $495\ \mu$ .<sup>1</sup> The organ of Corti of man has been estimated to include about 10,000 rods, and double that number of hair-cells and of auditory nerve-fibres.

We have up to this point traced the path of sonorous vibrations through the external and middle ear, through the ossicles to the fenestra ovalis, and thus to the fluid surrounding the terminal apparatus. We have now to examine the mechanism by which such vibrations may be supposed to effect auditory excitations.

According to the theory of Helmholtz, the stretched radial fibres of the basilar membrane behave towards the vibrations of the endolymph as the strings of a piano towards vibrations of the air caused by different notes. Different strings in the piano vibrate with different notes, particular strings respond to particular notes, and, in analogy with this known relation, it is supposed that different fibres in the basilar membrane vibrate with different notes, that particular fibres respond to particular notes. It is to be supposed that such responsive vibrations, although of infinitesimal magnitude, extend to the particular hair-cells resting upon the membrane, and give rise to excitations, which, conducted along nerve-fibres to the brain, produce

<sup>1</sup> These are v. Hensen's figures. According to Retzius, the inequality of breadth is much smaller, viz. 0.36 mm. at apex, 0.21 mm. at

different auditory sensations. On this theory it is supposed that the basilar membrane and hair-cells of Corti constitute an apparatus by which sound-vibrations are analysed and differentiated at the periphery; and as a supplementary feature of this theory, partly based upon the increasing breadth of the basilar membrane from below upwards, partly upon direct experiments upon dogs, it has been stated that low notes set into sympathetic vibration the upper fibres, high notes the lower fibres, of the basilar membrane. This theory has received support from experiments on dogs, and from the observations of von Hensen on the 'auditory' sacs of crustacea. These organs contain, besides otoliths, epithelial cells bearing hair-like processes, ranging from 140 to 720  $\mu$  in length. Different notes were found to set in motion different groups of hairs, but loud notes were found to set in movement all the hairs. The experiments on dogs are not very numerous or convincing; it is difficult to limit a lesion of the cochlea to its base or to its apex, and no less difficult to determine whether a dog has suffered in his appreciation of high and low notes; to be at all convincing, a dog who previously howled in response to high and low pitched music alike, should after destruction of the top of the cochlea respond only to high-pitched music, or, after destruction of the bottom of the cochlea, only to low-pitched music: this desideratum has not been fulfilled.

Rutherford, in 1886, proposed an alternative theory, in which the mechanism is compared with that of the telephone. He supposed that every cell of Corti is impressed by every audible vibration, and that the sound-wave transmitted by the fluid of the scala tympani through Reissner's membrane to the canalis cochleæ impresses the membrana tectoria and sets in vibration all the hairs of Corti's cells. He considers that through the intervention of these cells the vibrations are translated into nerve-impulses, which correspond with them in frequency, amplitude, and form, 'just as in a telephone the sound-vibrations are translated by the iron plate and magnet into electrical movements which correspond to those of the sound received.' He objects to the Helmholtz-Hensen theory, that a simple radial structure of the basilar membrane is by no means constant in all animals, that its increasing breadth from base to apex has been greatly over-estimated by Hensen, and that the probable consistence of the parts is not such as to suggest the occurrence of sympathetic vibration. According to Rutherford, complex vibrations are not



analysed at the cochlea, but translated into corresponding complex nerve impulses and sensations.

Hermann gives an experiment which is very unfavourable to the theory of consonation and peripheral sound analysis :—Two simultaneous tones,  $a_1$  (880) and  $c_{11}$  (1056), gave by the ear the beat-tone F (176) ; but it was not found possible to produce any such beat-tone in a distant resonator which was perfectly responsive to F as a simple tone. Two sounds, producing the usual beats when heard through one or both ears, can have a similar effect when each sound is heard exclusively through one ear. This, if correct, is significant of a cerebral fusion of stimuli from the two sides, and analogous with the binocular colour-mixture alluded to on p. 462.

A full consideration of the theories of auditory mechanism and of their respective merits would lead us too far ; we may remark in conclusion of this dubious subject, that the growth of evidence, so far from strengthening, has weakened the consonation theory. We may regard the basilar membrane as a long narrow drumhead, repeating the complex vibrations of the membrana tympani, and vibrating in its entire area to all sounds—although more or less in some parts than in others—giving what we may designate as acoustic pressure-patterns between the membrana tectoria and the subjacent field of hair-cells. In place of an analysis by consonation of particular radial fibres, it may be imagined that varying combinations of sound give varying pressure-patterns, comparable to the varying retinal images of external objects (w.)

There remains for us to trace the excitations made in one or other of the above ways, from their origin in the organ of Corti to their terminus in the cerebral cortex. The cochlear nerve-fibres are anatomically traceable to the spinal bulb by the dorsal root of the eighth nerve to the principal auditory nucleus and through the accessory auditory nucleus. From this station they doubtless proceed to the cortex cerebri, though along what precise paths, and to what precise region, it is not possible to say with assurance. It is, therefore, with considerable reserve that statements may be quoted to the effect that auditory channels pass to the corpus geniculatum posterum (testis) of the opposite side, and thence to the cortex of the first temporal gyrus.

Estimates of pitch and of intensity of sound are the effect of

cerebral comparisons with previous experience. The estimate of direction is mainly formed by inference from the differing intensities of the acoustic impressions on the two sides; a blindfolded person with the head kept stationary can easily localise correctly the direction of a sound produced to the right or left of the auditory field, but is very liable to misjudge the direction of sound directly in front or directly behind; he will easily localise correctly by moving the head to one side or the other.

The range of vibration-frequency within which tones are audible as such is from about 30 to 30,000 per second; but individual variations of auditory sensibility are considerable—of many persons the highest audible limit is 20,000, of others 15,000, or even less. Preyer gives as the high limit 40,000, and as the low limit 16. According to Zwaardemacher the high limit normally falls with advancing age from  $e_{vii}$  (about 40,000) at 10 years of age to  $b_{vi}$  (about 30,000) at 50 years of age. According to Exner, two sounds can be recognised to be distinct if the interval between them is not less than 0.002 sec. According to Schafhäutl, a person of acute hearing can detect the sound made by a cork ball weighing 0.001 gm. falling from a height of 1 mm. on a glass plate 91 mm. distant from the ear; this may be regarded as the ‘minimum audible sound.’

#### TASTE, SMELL, TOUCH, AND CUTANEOUS SENSATION

Taste and smell, although included under the classical five senses, and playing an important part in the selection of food-stuffs, are from an experimental point of view of very subordinate interest.

The principal nerve of taste is the glosso-pharyngeal, which supplies the posterior part of the tongue, *i.e.* that portion of the buccal surface which most contributes to taste. Two other nerves also take part in taste, inasmuch as their integrity seems to be necessary to the process; these are the so-called gustatory branch of the 5th, which is a common sensory nerve, and the chorda tympani of the 7th, the mode of action of which upon taste is, however, imperfectly understood. Sweets and bitters are best tasted when applied to the back of the tongue; salts, acids, and alkalies can be equally well ‘tasted’ by its lateral or anterior portions. There is a plant—*Gymnema sylvestre*—the leaves of which, when chewed, altogether abolish sweet and bitter taste.



Smell—‘the ancestral sense,’ inasmuch as the primitive brain is little else than an olfactory ganglion—is in man a comparatively insignificant mode of sensation and more important in its contribution to ‘taste’ than to smell proper. Taste is in reality in large measure effected by aroma. If smell is lost, or temporarily suppressed by filling the nostrils with fluid, very little discrimination by taste alone is left to us : most of our ‘tasting’ is by nose.

The nerve-ends by which taste-excitations are considered to be received are the *taste-bulbs*, which are most prominently distributed along the side of the trenches round the circumvallate papillæ, and which, in the rabbit, occupy the parallel trenches of two oval patches at the postero-lateral parts of the tongue. Each taste-bulb is an oval body formed of long fusiform cells, arranged in cortical and medullary groups ; the latter, to one pole of which nerve-fibres have been traced, project through an opening left at the opposite pole between the cortical cells.

The epithelium of that portion of the nasal mucous membrane to which the olfactory nerve-fibres are distributed has been termed the olfactory epithelium. It is composed of non-ciliated columnar cells, among which are scattered narrow, rod-like ‘olfactory cells’ with large oval nuclei.

**Cutaneous sensibility** includes common sensation, and is sometimes indiscriminately classed with a special sense—that of **touch** ; it is by means of cutaneous impressions that we are made conscious of the consistence, texture, and temperature of external objects, and it is partly by cutaneous sensation, partly by ‘muscular sensation,’ that we ascertain and compare their weight. As tactile organs, the skin of the palm of the hand and that of the sole of the foot play a most important part in our guidance : the varying pressures of the body on the plantar surface guide equilibration and locomotion ; we touch or feel objects by means of the hand and finger-tips, and we consciously utilise the sensations so acquired in our knowledge of objects and in our skilled movements.

The two most noteworthy subjects in connection with cutaneous sensibility are :—(1) its differences at different parts as ascertained by Weber’s compass method ; (2) the question of specific ‘heat nerves’ and ‘cold nerves,’ as investigated by Blix and by Goldscheider.

The nerve-ends to which tactile function is attributed are the touch-corpuscles of Wagner ; these, in the specially tactile area, *e.g.* the finger-tips, are set in ranks occupying rows of papillæ beneath the edges of the skin, forming thus what may be collectively regarded as a tactile organ. Each touch-corpuscle is a small oval glomerulus 50 to 200  $\mu$  long, formed of the branched and tangled ends of a medullated nerve-fibre, held together and surrounded by connective tissue. They are unequally distributed over the surface of the body, being numerous and crowded at the tips of the fingers, few and scattered on the arms, legs, and trunk, and our discriminative tactile power by means of palpation is in evident relation with this inequality of distribution. We can 'feel' things better with the finger-tips than with any other part of the body, although these have by no means the most 'sensitive' skin. Tactile sensation, although less obviously a special sense than vision or hearing, is different from common sensation, but unless we are on our guard there is risk of confusion, owing to the ambiguity of such familiar terms as 'feel' and 'sensitive.' We feel the pulse with the finger-tips, but to feel whether it rains or not we hold out the hand with the palm downwards ; if the palm of one hand and the back of the other are exposed together to a gentle drizzle, the droplets may be distinctly felt on the latter, while they are quite imperceptible on the former. We have already encountered a somewhat analogous distinction in the case of the retina ; the yellow spot, the region of most distinct vision, is not the most sensitive part of the retina (p. 459).

A measure of tactile discriminative power is obtained by finding the greatest distance at which two points of a pair of compasses applied to the skin are felt single, or the smallest distance at which they are felt double. Tested by this method (Weber's) the following are some average values in millimeters :

Side and dorsum of tongue . . . . .	9	Forearm (front) . . . . .	15
Tip of tongue . . . . .	1	Forearm (back) . . . . .	45
Palmar surface of hand . . . . .	8	Neck, back, arm, thigh . . . . .	50 to 70
Palmar surface of hand, terminal phalanx . . . . .	2	Cheek . . . . .	15
Palmar surface of hand, middle phalanx . . . . .	4	Lip (white) . . . . .	9
Palmar surface of hand, proximal phalanx . . . . .	5	Lip (red) . . . . .	4.5
Dorsal surface of hand . . . . .	30	Lip, inner surface . . . . .	20
		Forehead . . . . .	22.6
		Occiput . . . . .	27.1



By minute investigation of the skin, supplemented by subjective introspection, some remarkable conclusions have been arrived at, viz. that the skin contains many kinds of nerve-fibres and end-organs, separately subserving impressions of pressure, of pain, and of temperature. Of these investigations, the most noteworthy is that of Goldscheider, to the effect that different spots of skin are respectively excitable by heat and by cold, and that the nerve-fibres leading off from such 'cold' spots and 'heat' spots are specifically different from each other, as well as from the nerve-fibres leading off from the nerve-fibres leading off from 'pressure-spots,' and from 'pain-spots,' in accordance with the law of specific nervous energies.

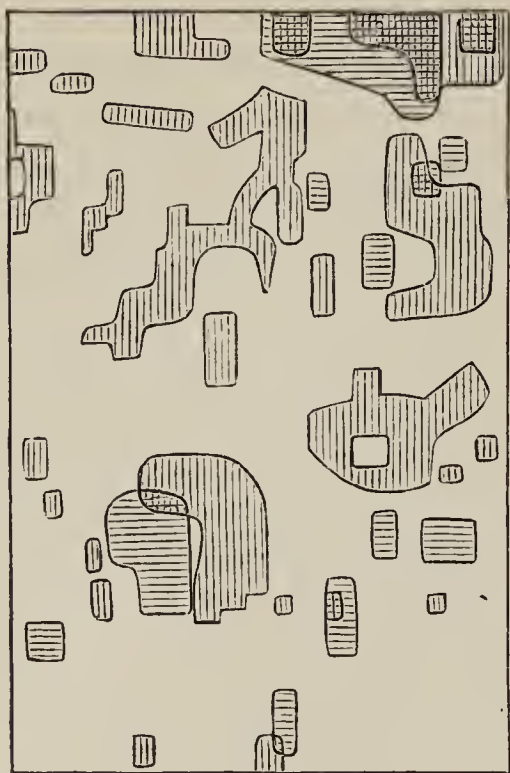


FIG. 262.

Goldscheider's maps of 'cold' spots' (vertical shading) and 'hot' spots (horizontal shading) superposed, and showing partial coincidence of 'hot' and 'cold' areas; anterior surface of thigh.

## CHAPTER XIV

## THE SPINAL CORD AND BULB

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## THE CRANIAL OR BULBAR NERVES

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**The spinal cord and bulb.** *Structure.*—The spinal cord consists anatomically of *white* matter and of *grey* matter; histologically, the white matter is chiefly composed of nerve-fibres: the grey matter is chiefly composed of nerve-cells and of nerve-fibrils; physiologically, the spinal cord is a *nerve-centre* by virtue of the cells of the grey matter, a *conductor* of nervous impulses by virtue of the nerve-fibres of the white matter, and in less degree by virtue of the nerve-fibrils of the grey matter.

Nerve-fibres in the white matter are medullated, vary greatly in diameter, and have no primitive sheath. Small fibres ( $5\ \mu$ ) predominate in the posterior columns, large fibres ( $15\ \mu$ ) in the cerebellar tract. Nerve-fibrils in the grey matter are for the most part the axis-cylinders of nerve-fibres and the branching



processes of nerve-cells. Nerve-cells in the grey matter vary greatly in size; 'large' cells (60 to 130  $\mu$  in diameter) are most

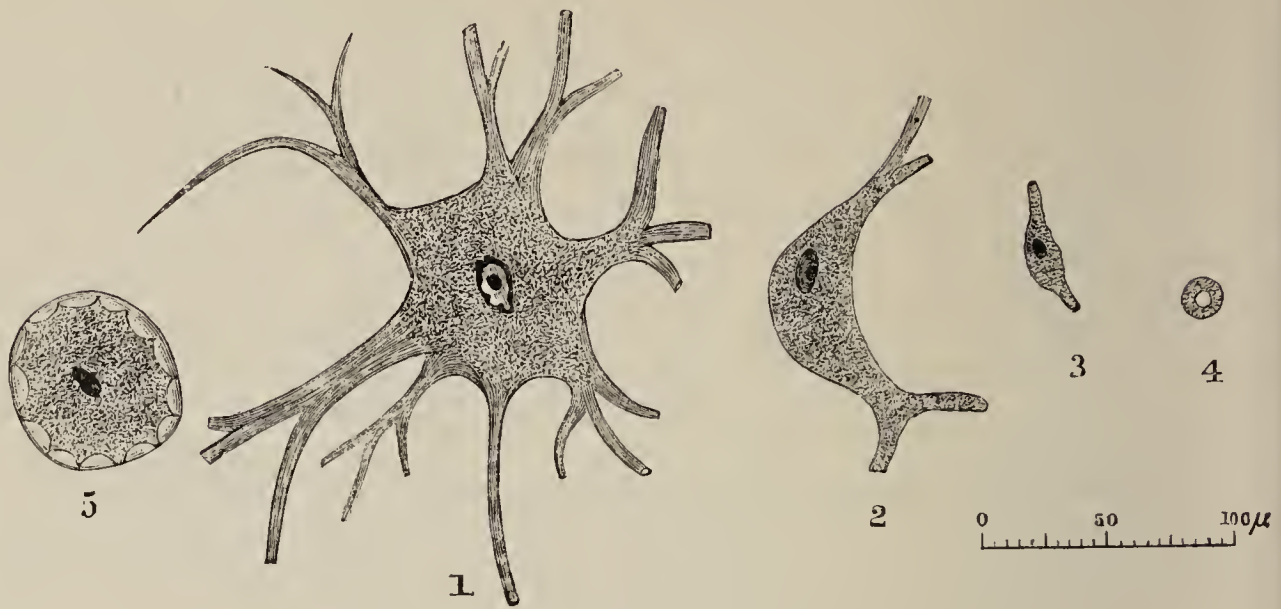


FIG. 263.

1. Cell of ant. cornu. 2. Cell of Clarke's column. 3. 'Solitary' cell of post. cornu. 4. Large nerve-fibre. 5. Ganglion-cell of posterior root. Drawn to the same scale, viz.  $\times 200$  diameters.

numerous in its anterior parts, and are connected with nerve-fibres of the anterior roots of the spinal nerves; 'small' scattered cells (20  $\mu$  in diameter) in the posterior parts of the grey matter are probably in relation with nerve-fibres of the posterior roots of the spinal nerves. The supporting tissue in which the true nervous elements are embedded is called the *neuroglia*; it is of a fine fibro-cellular structure resembling adenoid tissue, but, chiefly on account of its resistance to artificial digestion, it is classed with keratin, and supposed to have originated from the epiblast. It persists in greatest abundance in the *substantia gelatinosa*, which surrounds the central canal of the cord, and caps the tip of the posterior cornu. The central canal itself preserves through life a permanent vestige of its original formation by an infolding of the epiblast, in the form of a lining of ciliated epithelium. Nerve-cells and nerve-fibres are accumulated into longitudinal groups and strands, forming the various columns of the cord; strands of fibres form the *white columns*, longitudinal groups of cells form the *vesicular columns*. Grooves or fissures divide the white columns more or less superficially from each other into anterior, posterior, and lateral<sup>1</sup> columns. The relative situation of these several parts will be best understood by

<sup>1</sup> The term 'lateral' is here used in its original anatomical sense, and is comprehensive of the pyramidal and cerebellar tracts, as well as of the 'lateral columns' of neurologists.

studying them in a transverse section (fig. 264), and an idea of their longitudinal extension may be gathered from the diagram given in fig. 265. The white columns and the anterior vesicular groups extend through the whole length of the cord, and the latter are most prominent in the cervical and lumbar enlargements, the spinal centres from which the great nerves of the anterior and posterior extremities respectively take origin. The intermedio-lateral or lateral tract and the postero-median group of cells, known as Clarke's posterior column, are elongated islands rather than continuous columns of cells. They have their greatest development in the thoracic region, in which the

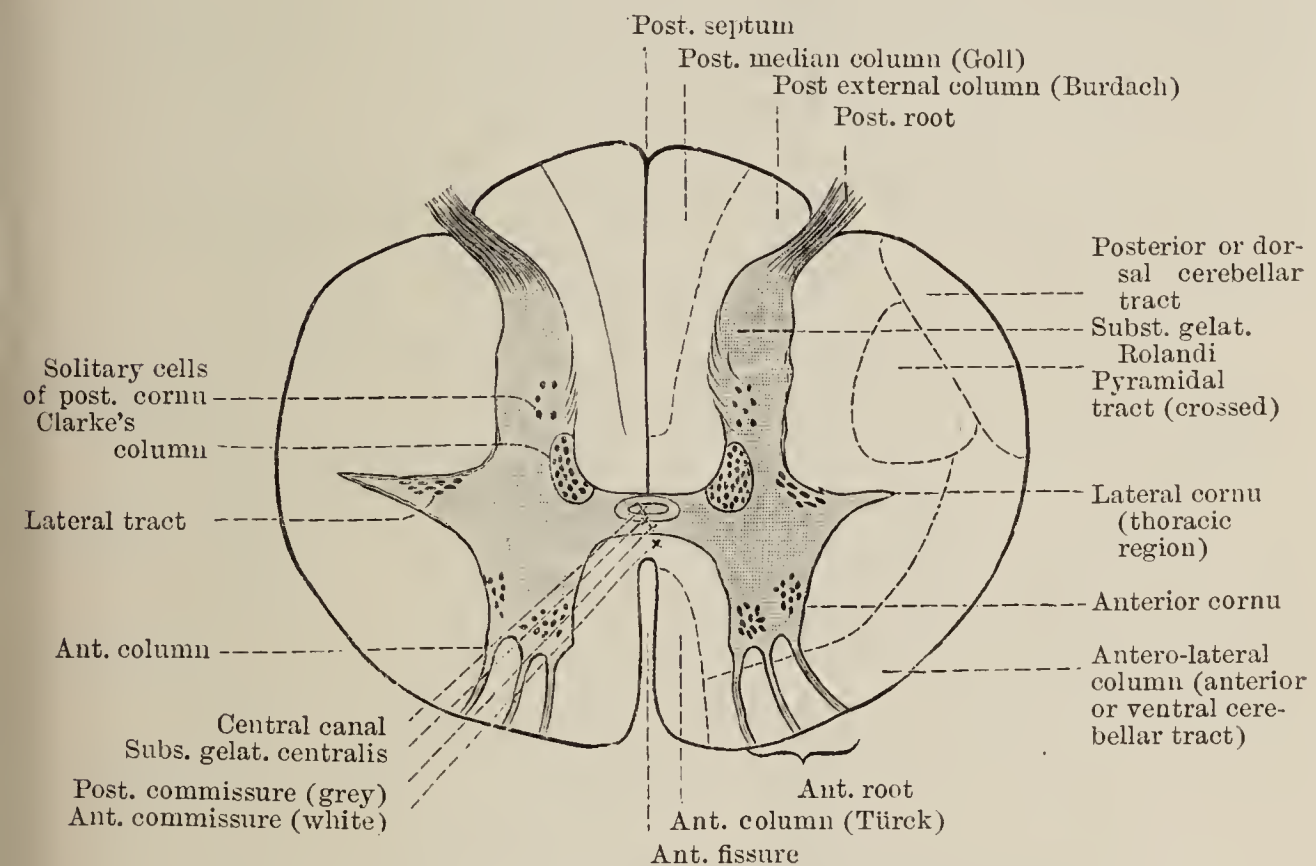


FIG. 264.—DIAGRAMMATIC TRANSVERSE SECTION OF THE SPINAL CORD,  $\times 6$ , on a level with the eighth thoracic nerve. (After Schwalbe.)

anterior columns of cells are most scanty ; in this region Clarke's column forms what is known as the *dorsal nucleus*. Similarly situated islands of cells are found in the cervical and in the sacral regions, known as the cervical nucleus and as Stilling's sacral nucleus. In the upper cervical region the lateral nucleus forms the nucleus of origin of the spinal accessory nerve ; in the bulb it is represented by the antero-lateral nucleus (=the chief vasomotor centre?).

At the bulb, in consequence of the opening out of the central canal into the fourth ventricle, and of the pyramidal decussation, the columns of the cord diverge and interlace, and its central grey



axis is broken up into scattered nuclei of grey matter forming the medullary reticulum and insular nuclei; these last coming close to the posterior aspect of the bulb. Among them are to be distinguished the vagus and hypoglossal nuclei, the gracile and cuneate nuclei, and the antero-lateral nucleus.

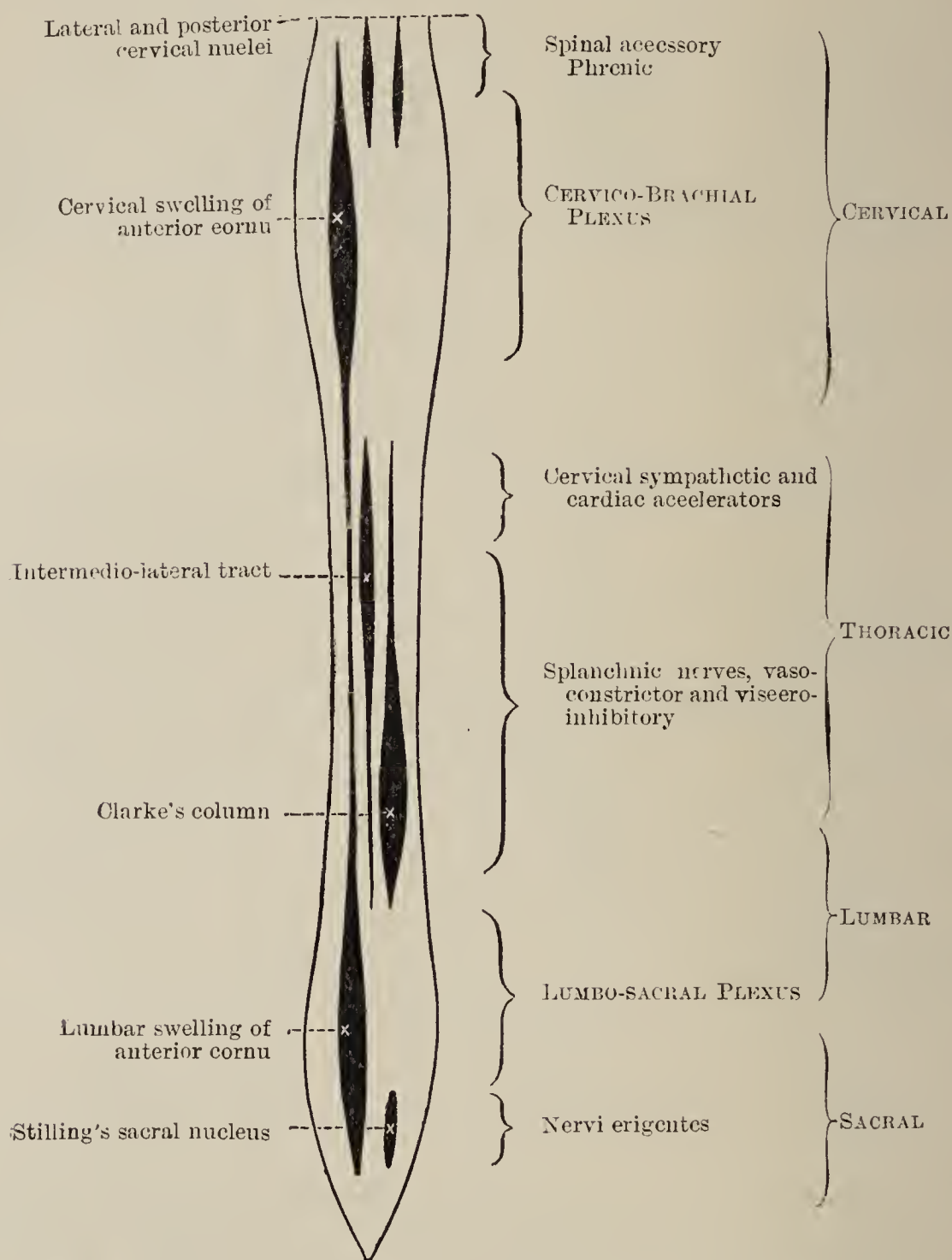


FIG. 265.—SIDE-VIEW OF AN IDEAL LONGITUDINAL SECTION OF THE SPINAL CORD.

To illustrate the distribution of the vesicular columns. The levels of origin of the principal nerves are indicated by brackets.

**Functions.**—That the spinal cord as a whole is a nervous *centre* admits of very simple proof, on the lower and on the higher animals, and on man. A decapitated frog having lost brain and bulb, reacts by movements when the skin is pinched, and no longer so reacts when the spinal cord is destroyed. A man whose spinal cord is cut, say in the dorsal region, by

disease or by mechanical injury, reacts by movements of his lower limbs when the soles of his feet are touched, and he bears witness to the fact that he feels nothing, that he is not conscious of any impression, *i.e.* that the reaction of the spinal cord is carried out without sensation. Such reactions are instances of reflex action in the original and restricted sense of the expression; they are simple, immediate, fatal and unfelt responses to unfelt peripheral stimuli.

Injury of definite transverse slices of the cord interferes with these reactions in definite transverse slices of the body; disease or injury of the cervical region abolishes reactions to and from the upper limbs, while reactions to and from the lower limbs may be unaffected; disease or injury of the lumbar region

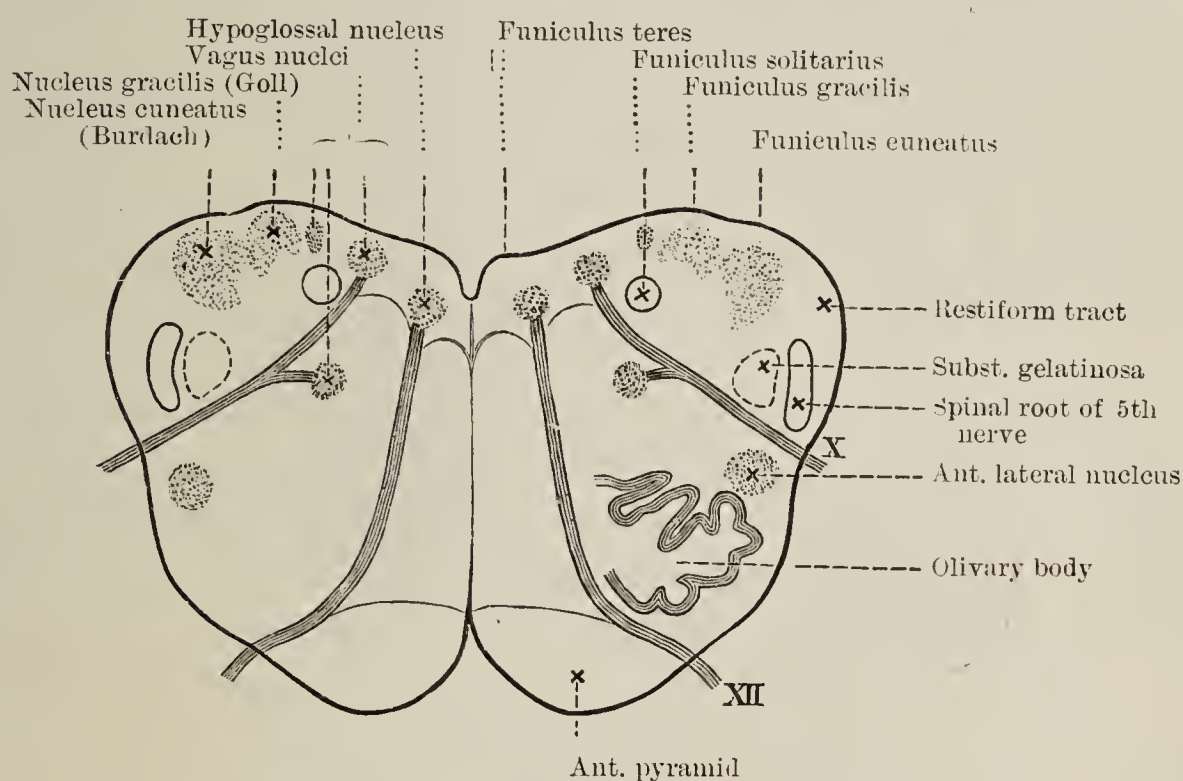


FIG. 266.—DIAGRAMMATIC TRANSVERSE SECTION OF THE SPINAL BULB,  $\times 3$ ,

at about the middle of the olivary body, to illustrate the principal nuclei and tracts at that level. (After Schwalbe.)

abolishes reactions to and from the lower limbs, while reactions to and from the upper limbs remain intact. This segmentation of spinal function applies to all vertebrate animals from man downwards; it is illustrated clinically by the various symptoms that characterise inflammation of the cord, when such inflammation is mainly lumbar, dorsal, or cervical, or when it progresses from one region to another: and it may at once be experimentally recognised on a decapitated frog by dividing the cord in the mid-dorsal region, when it will be found that reflex action can be obtained from leg to leg, or from arm to arm, but not from leg to arm, or from arm to leg.



The grey matter of the cord is thus to be regarded as a series of *segmental centres* fused together into a continuous longitudinal mass, each centre giving off a pair of spinal nerves to the two symmetrical halves of a vertebral segment. In correspondence with this view we may expect to find that the amount of grey matter is greatest on a level with greatest development of the vertebral segments, *i.e.* on a level with the limbs. This, indeed, is the case to such an extent as to give considerable swellings of the grey matter with which the large

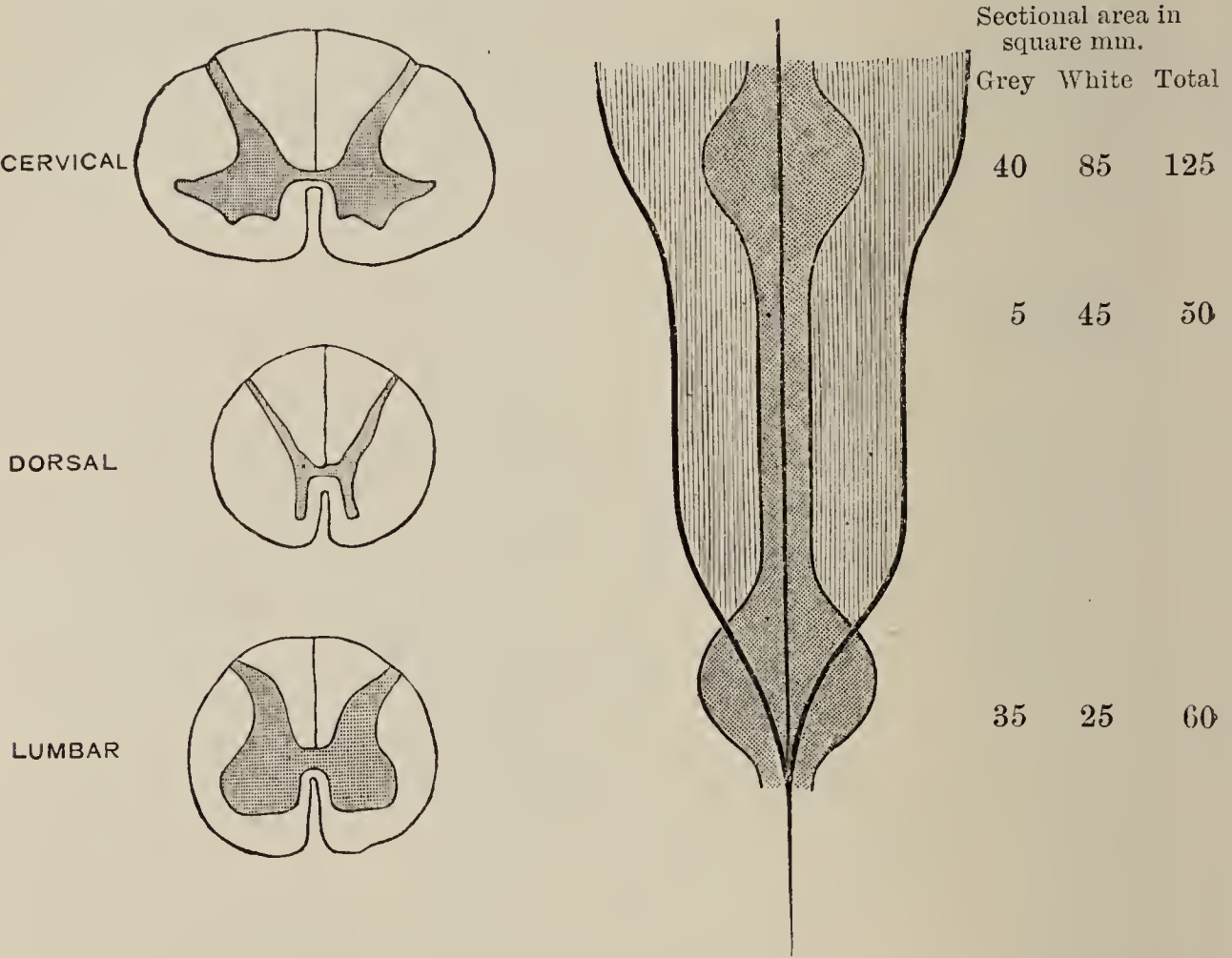


FIG. 267.—OUTLINE-SKETCH OF THREE SECTIONS ( $\times 3$ ).

Taken from the cervical, thoracic, and lumbar regions of spinal cord (human).

FIG. 268.—DIAGRAM-SKETCH BASED ON STILLÉ'S MEASUREMENTS.

To illustrate the area of white and of grey matter at different levels.

nerves of the upper and lower limbs are connected; and even in the cords of animals that show no such swellings, we find that the number of cells is greatest at points whence large nerves take origin.

The spinal cord is also a *conductor* of nerve-impulses. The white matter of the cord is in this respect to be regarded as a collection of nerve-fibres establishing connection between the brain and the grey matter of the spinal cord; the amount of white matter diminishes from above downwards progressively

with the smaller number of spinal cells that remain to be reached by such fibres; thus, at its upper part the white matter contains fibres sufficiently numerous to supply all the cells below, at a lower level it still contains fibres sufficiently numerous to supply all the cells below this level, but the number of cells in the latter case is smaller than in the former, and the number of fibres thus diminishes downwards. It is on this account, and not because they lose fibres into each pair of spinal nerves, that the white columns taper downwards.

There are reasons for accepting it as highly probable that there are no direct fibres between the cortex cerebri and the periphery (sensory or motor), *i.e.* no fibres to or from the cortex that have not a spinal cell in their course. Grounds for these statements are furnished—(1) by actual enumeration of fibres and of cells throughout the spinal cord of a small animal—a laborious task, which has been undertaken in the case of the frog only; (2) by tracts of degeneration, *i.e.* ‘descending’ degeneration of the pyramidal tracts tapers downwards, and does not extend beyond anterior nerve-cells into motor nerves; ‘ascending’ degeneration of Goll’s column is not traceable beyond the bulb (nucleus gracilis).

**Nerve-roots.**—Afferent and efferent fibres are intermingled in mixed nerves; in the roots of the spinal nerves they are separated, the anterior roots being composed of efferent or motor fibres, the posterior roots being composed of afferent or sensory fibres. Section of the anterior roots causes paralysis of motion; section of the posterior roots causes paralysis of sensation; excitation of the central end of a divided anterior root gives no effect, excitation of its peripheral end causes muscular contraction; excitation of the central end of a divided posterior root causes reflex movements, excitation of its peripheral end has no effect. One of these four statements requires qualification—excitation of the peripheral end of a divided anterior root, if the test be applied after the ‘shock’ caused by the preparatory operation has passed off, will evoke reflex as well as direct movements; the anterior root, although mainly composed of motor fibres, contains a few sensory fibres derived from the posterior roots, and therefore termed recurrent sensory fibres. All these facts, inclusive of those dependent upon the existence of recurrent sensory fibres, were discovered by Magendie (1822). Charles Bell (1811), to whom they are commonly ascribed, observed motion on excitation of the anterior roots of a recently



killed animal, and inferred that the anterior or 'cerebral' roots are motor and sensory, and that the posterior or 'cerebellar' roots 'serve to govern vital actions.' A. Walker (1809) made the unfortunate guess that the anterior roots were sensory and the posterior motor.

Impulses along recurrent sensory fibres must run a very exceptional and often tortuous course ; the fibres do not, as was

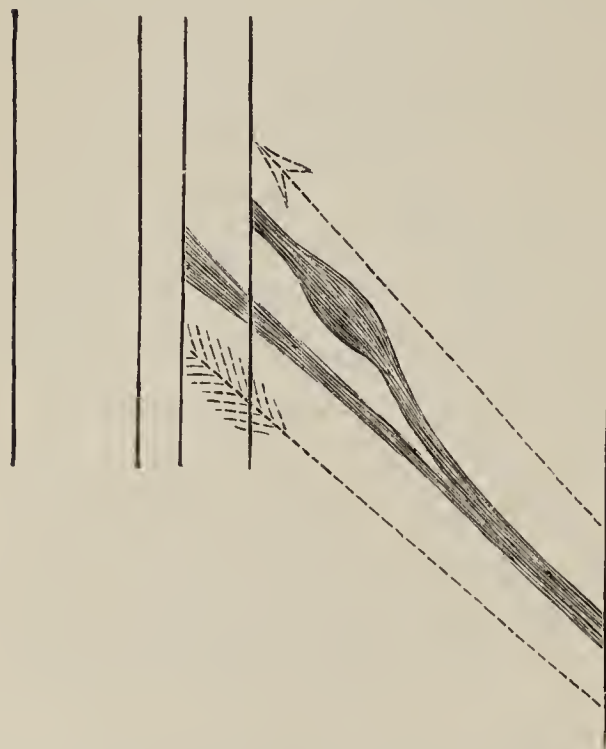


FIG. 269.—RECURRENT SENSORY FIBRES.

The probable course of impulses along them indicated by the dotted arrow ; their degeneration occurs 'against' the arrow, their trophic centre being the posterior root-ganglion.

first supposed by Magendie, turn back close to the junction of the roots, but they run in the nerve-trunks towards, and even into, the peripheral plexus of nerves, whence they return to the posterior roots (Bernard). It is probably owing to the excitation of recurrent sensory fibres that the excitation of the *peripheral*<sup>1</sup> ends of many nerves—motor as well as sensory (*e.g.* fifth and seventh)—is found to produce manifestations of pain, and that sensation, temporarily abolished by 'shock,' can reappear a day or two after nerve-section (*e.g.* of the median nerve) without it being necessary to invoke any

sudden reunion of the divided nerve.

The two pairs of nerve-roots connected with each spinal segment contain all the efferent and afferent fibres to and from the tissues—internal as well as external. All efferent fibres emerge by the anterior roots ; their most prominent variety are the motor fibres of voluntary muscle, but they also include vasomotor, visceromotor, pilomotor, and other fibres. All afferent fibres enter by the posterior root ; their most prominent variety are the sensory fibres of the skin, but they also include visceral and 'muscular' fibres. It has been supposed that the motor fibres contained in a given root are functionally associated, seeing that excitation of a root was found to set in motion muscles that habitually act together (Ferrier and Yeo), but an accidental muscular combination is difficult to distinguish from a true functional combination ; and (although, doubtless, functional association and anatomical contiguity must coincide to some extent) it is more probable that each

<sup>1</sup> The peripheral end of a divided nerve is the end attached to the periphery, the central end is that attached to the centre.

single root is an anatomical group, in accordance with spinal segmentation (Müller, Sherrington), rather than a functional group. The anatomical association of different sensory fibres in a given root is of considerable physiological and clinical significance; each spinal segment is a station with which are connected fibres from the skin, of which our sensation is most distinct, and fibres from the viscera, of which our sensation is most indistinct. Visceral impulses of sufficient intensity to arouse a spinal segment may cause tenderness or actual pain in a part of the skin served by the cutaneous sensory fibres connected with that segment; the seat of cutaneous tenderness may, therefore, be an index to the seat of visceral disease (Ross, Head).

It is most natural from a physiological standpoint to speak of nerve-cells as being in protoplasmic *continuity* with nerve-fibres; but, according to modern histologists (Golgi, Ramón y Cajal, Retzius), such continuity is in many instances absent, the relation (which applies to peripheral as well as to central cells) being one of *contiguity* between nerve-cell and a surrounding network of fine fibrils. An anterior cornual cell, *e.g.*, is continuous with an efferent nerve-fibre by an axis-cylinder process, but contiguous only with a pyramidal fibre by a surrounding ramification of fibrils. But this anatomical discontinuity does not involve functional discontinuity: a nerve-impulse along a pyramidal fibre can *influence* a 'motor' cell, and physiologically we may picture the process as taking place along a continuous strand of excitable protoplasm. With regard to the actual significance of the interruption, we may surmise: 1. That it plays the part of a 'nerve-valve,' permitting impulses to pass in one direction, while blocking them in the reverse direction. 2. That it is a determining factor as regards the trophic action of nerve-cell upon nerve-fibre; *e.g.* the anterior cornual cells (1) permit the passage of centrifugal impulses in the direction *a, c, e*, and obstruct that of centripetal impulses in the direction *e, c, a*; (2) have a trophic action upon efferent root-fibres at one pole *c, e*, but no such action upon pyramidal fibres at the other pole *a, c*, the nutrition of the latter being controlled by a Rolandic cell.



FIG. 270.

**Paths of motor and of sensory impulses in the spinal cord.**—A paralysis of motion or of sensation, caused by a lesion on one side of the brain, is on the side of the body opposite to the lesion. This connection, which is practically constant, proves that motor and sensory paths must cross the middle line. Where do these channels cross, and how far can we precisely define them? We shall find that this question can be answered with an approach to certainty as regards motor paths, but with far less assurance as regards sensory paths. Data from three different sources



contribute to our knowledge of the matter :—(1) Data obtained after experimental lesions upon animals, or accidental lesions upon man ; (2) data obtained by the study of tracts along which degeneration is traced some time after the occurrence of such lesions ; (3) data obtained by the study of the order of development of various tracts.

(1) **Experiments.**—The effect of complete transverse division of the cord is *paraplegia*, *i.e.* paralysis of voluntary motion and of sensation in all parts below the injury. The effect of hemisection, *i.e.* of transverse division of one-half of the cord, is paralysis of voluntary motion on the same side, with weakness on the opposite side, and paralysis of sensation on the opposite (?) side. Sensibility has been found exaggerated on the same side ; by other observers it has been found that the loss of sensibility is greatest on the same side. The effect of longitudinal division in the middle line is said to be an abolition of sensation on both sides, and a weakness of movement not amounting to complete paralysis. The effect of two transverse hemisections on opposite halves of the cord varies according to the interval between them ; if near together, two hemisections are equivalent to a complete transverse division ; if far apart, motility and sensibility are more or less preserved.

The generally quoted conclusions from these experiments are, that motor channels cross at the *bulb*, sensory channels in the *cord*. Both statements must be qualified, and we may say at once that, whereas precision will be added to the first statement, reservations to the second must be made almost amounting to contradiction. With regard to *motor* channels, while their major part cross at the decussation of the pyramids, a minor part (in man) cross in the spinal cord, and we shall see that these parts are picked out by degeneration, as well as recognisable by their tardy development, as the crossed and direct pyramidal tracts. But with regard to *sensory* channels the evidence is far more uncertain and unfinished. Clinical evidence shows that the sensory path crosses somewhere, but not where ; until recently the statement was repeated unchallenged, and by mere force of repetition acquired, perhaps, undue credit, to the effect that sensory channels cross in the cord. Now, however, there is an increasing disposition to abandon this view ; the recent experiments of Gotch and Horsley, and those of Mott, have yielded evidence in opposition—the electrical token of centripetal impulses has been found most pronounced on the *same* side

of the cord as excitation; monkeys after hemisection have been found with the less sensitive leg on the *same* side as the lesion, as was originally stated by Galen, 1,700 years ago, presumably from experiments on monkeys.<sup>1</sup> To such opposition evidence must also be added the indirect testimony of degeneration—‘ascending’ degenerations above a hemisection are most marked on the same side of the cord, and degenerations between the bulbar nuclei and the cerebral cortex cross the middle line at the decussation of the fillet. It appears then that sensory as well as motor channels have their major decussation in the bulb, their minor decussation in the cord; in which case we must admit that sensory impulses from each side of the body can pass up both sides of the cord, but most on the same side.

The older experiments above alluded to originate from the Paris school, and are due chiefly to Brown-Séquard. More detailed experiments on the same subject have been made in Germany by Schiff, Türck, Stilling, Müller, and by Ludwig’s pupils. The main result of these last was to the effect that the lateral columns of the cord are the chief channels of motion and of sensation, and that the anterior columns take part in motor conduction; vasomotor, among other motor effects, were shown to have their passage in the lateral columns of the cord and in the anterior roots of the spinal nerves; after hemisection of the cord the vessels were found to be dilated on the same side below the injury, but excitation of the cord above the hemisection causes vaso-constriction on *both* sides (Nicolaides).

Attempts have been made to further define what kinds of motion and of sensation pass along the various columns. Separate tracts have been assigned to *voluntary*, as distinguished from automatic or reflex motion, and although we must not hastily admit special and exclusive functions of particular strands, we may recognise the pyramidal tracts as pre-eminently the channel of ‘voluntary’ action; they connect the Rolandic region of the brain with the anterior-cornua of the cord, and are more voluminous in man than in any other of the mammalia. Separate tracts have been assigned to various modes of sensation. ‘Tac-

<sup>1</sup> ‘ . . . Moreover, you have seen that transverse incisions of the whole cord deprive all parts of the body situated below of sensibility and of movement. . . . And you have seen in dissections that transverse incisions of the cord which stop at its centre do not paralyse all the inferior parts, but only the parts situated directly below the incision—on the right, when the right side of the cord has been cut; on the left, when it is the other side’ (Galen, ‘X. Of the Parts Affected,’ lib. iii. cap. xiv.; Daremberg’s transl. II. p. 579).



tile sensibility' has been billeted upon the posterior columns, 'sensations of pain' have been considered to be received through the grey matter, common sensation through the lateral columns, thermic sense and muscular sense through the posterior columns. All these opinions are mainly based upon the shifting ground of subjective interpretations, and are therefore most doubtful and inconclusive. Signs of sensation are difficult to read in animals, signs of lost sensations are even more equivocal, and observations as to whether different kinds of sensation are preserved, or lost, or altered, are particularly liable to be guided by the expectations of the observer. In man, the balance of evidence seems to be in favour of Schiff's and of Vulpian's conclusion, that the grey matter is the chief conductor of sensory impressions. In syringomyelia (*i.e.* dilatation of the central canal and degeneration of the grey matter) there is, among other symptoms, loss of sensibility to thermic and to normally painful stimuli, but not to tactile stimuli.

*Direct excitability of the spinal cord.*—It might be expected that direct excitation of various tracts of the cord should be capable of yielding information concerning the different paths followed by nervous impulses. But this has not been found to be the case; the exact localisation of experimental stimuli is extremely difficult to accomplish on the intact cord, and to artificially isolate longitudinal tracts with the knife depresses excitability to such an extent that it may then with difficulty be manifested, even with strong stimuli. It is for these reasons that diametrically opposed answers have been given to the question, 'Is the spinal cord excitable by direct experimental stimuli?' Van Deen, Schiff, and Chauveau, answered 'No.' Vulpian, Fick, and Dittmar, answered 'Yes.' In the attempt to form our opinion from such conflicting statements of fact, we must bear in mind (1) that the excitability of the cord is easily depressed by the shock of the operation necessary to expose it; (2) that the cord is of complex structure, being composed of (*a*) nerve-fibres of the anterior and posterior roots; (*b*) nerve-fibres of the white columns; (*c*) grey matter; and the question must therefore be put separately to each of these three constituents. The nerve-fibres of the anterior and posterior roots pass obliquely through the white columns for a short distance above and below the level at which they emerge; the excitability of these root-fibres is admitted by all observers. By those who deny the direct excitability of the white columns, the muscular contractions which have been observed to ensue upon

mechanical or electrical stimulation of the columns, are attributed to stimulation of these root-fibres. The nerve-fibres of the white columns are admitted to give passage to nerve-impulses between brain and cord; it is difficult to admit their inexcitability to direct stimulation, and it is very probable that their frequent failure to give evidence of direct excitability has been due to shock. The evidence required to prove the negative proposition is in this case far greater than that required to prove the opposite positive proposition.

An experiment of Fick's, which has been repeated by other experimenters, furnishes such evidence with regard to motor columns. By longitudinal incisions he isolated long strips of white matter, and in favourable cases obtained contractions as the result of excitations applied far from the point of division. Under these circumstances the stimulus was localised to the columns, and could not affect root-fibres, which were cut. And in the experiments of Gotch and Horsley (p. 397), electrical effects were produced by excitation of the various columns above and below. We may therefore admit the excitability of the white columns as proved, though it has been difficult of proof.

To test the direct excitability of the grey matter is a still more difficult task. That it can transmit motor and sensory impulses is admitted on all hands. That it is excitable by direct experimental stimulation cannot be asserted, fact in hand, for it occupies a central situation, surrounded by nerve-fibres in close proximity. The fact is not proven. We may, however, remember that the grey matter of the cerebral cortex, which is accessible, was long supposed to be inexcitable by direct stimuli, but is now generally admitted to possess direct excitability; the fact is, therefore, probable.

Excitability by experimental stimulation is not to be confounded with indirect or functional excitability evoked by stimuli that reach the cord by the natural channels. In this sense it is obvious that the cord (both white and grey matter) is excitable. A classical experiment of Stenson's demonstrates how rapidly the functional excitability of the grey matter is lost when the blood-supply is arrested. The abdominal aorta in the rabbit can be completely compressed so as to arrest circulation in the lumbar part of the cord; if this arrest be maintained for two or three minutes, the posterior extremities become completely paralysed as regards sensation, voluntary motion, and reflex motion; the nerves and muscles are not paralysed, but the grey matter of the



cord is rendered inexcitable ; if the arrest is much prolonged (an hour or more) the cells may not recover at all, and the fibres in connection with them will then degenerate. The experiment furnishes one among other proofs that motor and sensory channels all pass to and from the brain *via* a station of grey matter in the cord, and that there are no direct channels of white matter in either direction between the brain and the periphery. Arrest of circulation for so brief a period, though sufficient to abolish the excitability of nerve-cells, is not sufficient to abolish the conductivity of nerve-fibres.

The same fact is demonstrable on decapitated animals. Excitation applied by means of electrodes inserted into the vertebral canal a minute or two after decapitation gives contractions of the upper extremities (by direct excitation of motor nerve-fibres), but no contraction of the lower extremities, because conduction is interrupted by inexcitable grey matter (w.)

(2) **Degenerations.**—After transverse division of the cord, either experimentally upon animals or accidentally upon man—*e.g.* in the mid-dorsal region—degenerations occur along certain tracts above and below the seat of division, in consequence of the separation of fibres from their ‘trophic’ cells ; the former—called ‘ascending’—are found in the column of Goll and in the cerebellar tracts ; the latter—called ‘descending’—are found in the column of Türk and in the lateral column. A complete transverse division of the cord gives, of course, no information whether or no degenerated tracts cross the middle line ; hemisections, or unilateral lesions, are necessary to supply such information. As regards descending degeneration, however, abundant data are furnished by cases of cerebral hemiplegia, where, in consequence of a brain lesion, a definite tract of fibres degenerates, and can be traced in the brain, in the bulb, and in the cord. In the latter, the degeneration consequent upon a lesion on one side of the brain occupies two definite situations in a transverse section of the cord ; the larger tract of degeneration (crossed pyramidal tract) occupies the lateral column of the side opposite to that of the cerebral lesion, the smaller tract (direct pyramidal tract) occupies the anterior column of the same side as that of the lesion. The fibres composing this smaller tract are believed to cross the middle line along the whole length of the cord, to be distributed to the opposite side of the body. This coincides with the results of the experiments given above, to the effect that motor channels cross in major part at the decussation of the

pyramids, in minor part throughout the spinal cord. The degenerated tracts taper downwards in both cases, the crossed tract being traceable farther down the cord than the direct tract; the degeneration does not extend to the anterior roots, *i.e.* it is limited by the cells of the anterior cornua.

Ascending degeneration after hemisection of the cord, *e.g.* in the dorsal region, takes place on the same side as the injury, in the postero-median column up to the nucleus of the funiculus gracilis, and in the cerebellar tracts. It may also occur in consequence of destruction of spinal ganglia, or of division of roots between ganglia and cord; in this case the degeneration is limited to the posterior columns and absent from the cerebellar tracts—facts which are taken to signify that ‘ascending’ fibres of the latter are connected with cells in the cord below (cells of Clarke’s

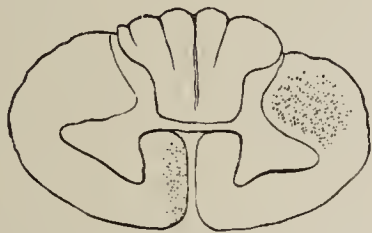


FIG. 271.—DEGENERATION OF THE SPINAL CORD.

Secondary to a lesion of the right hemisphere; *right* direct pyramidal tract, *left* crossed pyramidal tract (Mott). (See also fig. 284.)

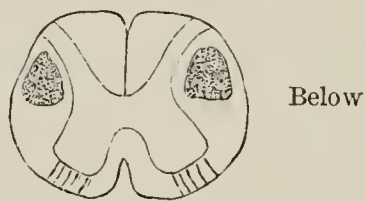
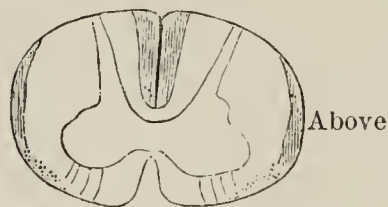


FIG. 272. - TRACTS OF DEGENERATION ABOVE AND BELOW A TRANSVERSE DIVISION OF THE CORD.

column), while fibres of the posterior column are in uninterrupted continuity with posterior root-fibres and ganglia. These fibres do not, however, pursue an unbroken course to the cortex; the degenerated tract ends at the spinal bulb in connection with the cells of grey matter, of the gracilis and cuneate nuclei, which are the bulbar representative of the grey matter forming Clarke’s column. From these bulbar nuclei fibres are traceable (1) to the opposite side of the cerebellum *via* the superior pyramidal decussation and restiform body; (2) to the opposite side of the cortex cerebri *via* the fillet decussation and internal capsule. The posterior or dorsal cerebellar tract is traceable through the restiform body into the cerebellum; the anterior or ventral cerebellar tract extends higher and passes to the cerebellum through its superior peduncle (Mott). Ascending degeneration of Goll’s column tapers upwards, a fact that is referable to the connection of its fibres



with the cells of Clarke's column; degeneration of the posterior roots, followed up the cord, is at first postero-lateral, and higher up postero-mesial in situation, so that in a section across the cervical region, fibres derived from lower roots are situated on the mesial side of fibres derived from the upper roots.

The above-described are the main tracts along which systematic degenerations have been followed down and up the cord; we may, however, add, without entering upon comment or discussion—(1) that scattered 'ascending' as well as 'descending' fibres, in more or less abundance, have been found degenerated in other situations—in the antero-lateral and in the posterior columns; (2) that a small patch of fibres in the posterior column (referred to by pathologists under the name of 'comma tract') undergoes descending degeneration after section of the posterior roots; (3) that ascending and descending degenerations taken together do not include the entire area of the white columns, but leave a considerable remainder, especially in the immediate vicinity of the grey matter, the fibres of which suffer no degeneration, and are therefore considered to be commissural between cells at different levels; (4) that in cases of locomotor ataxy a well-marked area of degeneration occupies the postero-external columns (fig. 264) and the postero-median columns at higher levels.

It should, moreover, be expressly stated that tracts of degeneration do not constitute direct evidence that such tracts are, normally, functional channels in the direction of degeneration. They indicate lines of trophic influence derived from 'centres' which have been destroyed, or from which the fibres have been separated. It is, however, considered probable that such lines of trophic influence coincide with lines of functional impulses—but we must recognise that the inference is not unimpeachable. As a matter of fact, we know that afferent nerves of the posterior root divided beyond the ganglion, undergo 'descending' degeneration. The terms are also open to the objection that, as in the case of nerves, the degeneration does not gradually ascend or descend, but that it simultaneously invades the entire length of a tract separated from its trophic centre.

(3) **Development.**—The various white tracts of the spinal cord are not developed simultaneously, but successively; the fibres composing them acquire their medullary sheaths at different dates, so that in embryos at various stages various tracts may be distinguished and separately followed. As regards this order

we may, without going into details, quote the main conclusions of Flechsig's investigations: the first apparent fibres are the peripheral or root-fibres connecting the spinal grey matter with the periphery—sensory, then motor; the next are the commissural fibres connecting grey matter at different levels; next, the centripetal tracts from cord to bulb and cerebellum; finally, the centrifugal tracts from cerebrum to cord. The most important point which has been thus brought out is, that the pyramidal tracts above described as degenerating in consequence of cerebral lesion are the last-comers, and that they appear simultaneously with the cortex cerebri from which they take their origin. This is corroborative evidence of the view that the pyramidal tracts are motor channels from the cortex. On man this motor or pyramidal system does not appear until birth or a few weeks later, before which it is not possible to distinguish a pyramidal tract by any of the methods used to demonstrate tracts of medullated fibres; these tracts are not well developed until the end of the first year, and even then are recognisable as the youngest of the spinal tracts, being composed of fine fibres ( $2\ \mu$ ), whereas in the adult state they are second only to the direct cerebellar tracts as regards the diameter of the majority of their fibres (5 to 10 to  $15\ \mu$ ). In the development of the pyramidal tract two further points are distinctly noteworthy. Firstly, the fact that its nerve-fibres go through three recognisable stages:—appearing as naked axis-cylinders (fifth month after birth); as fine medullated fibres (ninth month); finally, as coarse medullated fibres. Secondly, the probability, almost amounting to certainty, that the pyramidal tract, appearing as the last-comer in the cerebro-spinal axis, develops in a centrifugal manner, pushing and insinuating itself along the lines which it finally occupies. A third point should also be alluded to, viz. that the relative proportion between direct and crossed tracts is liable to vary; ordinarily, the direct is to the crossed as 1 is to 9, but the proportion may be occasionally 1 to 4 or 1 to 1, and, very exceptionally, the entire pyramidal mass may fail to cross, being continued as an abnormally large anterior column.

To sum up the considerations contained in the foregoing pages—experiment, clinical observation, the study of development and of degeneration, concur to testify that in man the *motor* channels from one side of the brain pass to the opposite side of the body *via* the lateral column of the opposite side and the anterior column of the same side; the former channel, called



the crossed pyramidal tract, is the larger, and crosses the middle line in the bulb; the latter channel, the direct pyramidal tract, is the smaller, and crosses the middle line in the spinal cord. As regards *sensory* channels, it is proved that these cross from one side of the body to the opposite side of the brain, but as to the precise locality of the crossing, evidence is conflicting. According to the older experiments, the chief crossing is in the cord; according to newer experiments, and to the collateral but not perfectly conclusive evidence of development and of ascending degeneration, the chief crossing is in the bulb, above the decussation of the pyramids.

**The spinal cord as a centre of reflex action.**—We have seen that the grey matter of the spinal cord constitutes a series of centres, and that the movements over which they preside are pre-eminently of a *reflex* character. Certain movements, however, of undoubted spinal mechanism stand out from this category, and are more commonly characterised as *automatic*; certain other movements, also of undoubted spinal mechanism, demonstrable only on the lower vertebrates, are manifestly appropriate to definite ends, and have the appearance of being volitional in character to such a degree that the expression *psychical* has even been used to characterise them. Yet these movements in no wise detract from the statement that the spinal cord is the typical centre of reflex actions in their purest form, namely, immediate, unchosen, fatal responses to peripheral excitations, without the intervention of consciousness and volition. The aimless movements of the limbs of the paraplegic patient are a glaring instance of this kind of action in its lowest form. A grade higher, and the reflex act is something more than an aimless spasm: it is a defensive act with animal self-preservation as its result, a character that may very generally be detected in spinal reflex acts. A grade higher still, and the spinal reflex act is so frequent and habitual as to *appear* to have become independent of peripheral excitations, and now it receives the name of an automatic action. Such actions are very evidently defensive and self-preservative as regards animal life. No sharp boundary-line can be drawn between the reflex and the automatic. As we have seen in Chapter VIII, the two expressions overlap, and are often used indifferently; but, contrasting extreme cases of each, we may say that the reflex act is occasional and its excitation definite, while the automatic act is habitual and its excitation indefinite.

Further analysis of the mode of action of the spinal cord as a centre can be best made on decapitated frogs. The following are the chief experiments and observations that may be repeated.

1. *Summation of stimuli*.—A series of comparatively weak stimuli is far more effectual in eliciting a spinal reflex action than a single stimulus of much greater strength. This may be demonstrated by applying induction shocks to the skin, or, better (in order to eliminate the possibility of the summation having its seat in the cutaneous nerve-endings), to the central end of a divided sciatic nerve. Very weak tetanisation is effectual, while much stronger single shocks give no effect at all; but even single shocks will become efficacious above a certain limit of strength. To use an analogy—a nail is driven home better by a series of comparatively light blows than by one single heavy blow.

2. *Diffusion of stimuli*.—If the skin of one limb be stimulated by induction shocks of minimal strength, the first muscular reaction occurs in that limb. If the strength of stimulation be increased, the opposite limb of the same segment also contracts; on further increased strength of stimulation, the other limbs enter into action. Thus, if the skin of the right inferior extremity be stimulated, the muscular responses occur—(1) In the right inferior extremity; (2) in the right and in the left inferior extremities; (3) in the inferior and in the superior extremities. That is to say, excitation diffuses in the grey matter of the spinal cord, first transversely, then longitudinally.

3. *Inhibition of stimuli*.—Two successive stimuli on the same spot summate. Two successive stimuli on different spots may either summate or interfere with each other. Two simultaneous stimuli on different spots more commonly interfere with each other. These statements are not to be taken as absolute, but only as expressing a general rule, to which there are many exceptions. The best experiment to demonstrate inhibition of stimuli in the spinal cord is made on the decapitated and brainless frog by means of chemical stimulation. The tip of one toe is stimulated by being dipped into weak acid ( $\text{H}_2\text{SO}_4$ , 2 per 1,000), and the interval between application of stimulus and retraction of the limb is measured. A crystal of common salt is applied to the upper part of the cord and the experiment is repeated; the reaction interval is found to be longer, or it may be necessary to use strong acid to get any reaction at all. This mode of testing the excitability of the cord by stimulating the skin with weak acid is known as Türk's method; the experiment



above described with salt is a modification of one by Setschenow, in which it is applied to the optic lobes in order to demonstrate their inhibitory action on the cord.

4. *Time of reflex action.*—Türck's method has been employed to determine the interval between stimulus and reaction, but is open to the objections that it necessarily includes the time during which the acid is soaking through the skin, and that the stimulus is in reality an increasing one, the exact moment of explosion of which it is impossible to determine. The real time of reflex action is best found by applying single induction shocks to the central end of a divided sciatic, and correcting the total time of reaction by the amount lost in transmission along nerves and in delay at the muscle. Measured thus, the true reflex time, *i.e.* that occupied by the elaboration of a stimulus in the spinal cord, is found to be only  $\cdot 01$  to  $\cdot 015$  second; whereas by Türck's method the interval between dipping and contraction may be anything between 5 and 30 seconds.

On mammalia the true reflex time of the spinal cord has been imperfectly studied. On an intact animal we cannot be certain that a movement, which is to all appearance reflex, is in reality effected through the spinal cord alone; it may be a reaction from the brain. This especially applies to man: the prick of a needle will cause a so-called reflex withdrawal of the hand, and the total interval between the two events will be above  $\frac{1}{10}$  second; but it is probable that this is a cerebral reaction involving sensation. Winking of the eyes in response to a stimulus applied to the conjunctiva, is the nearest approach to a true reflex act demonstrable on man; the organ of return of action being in this case the spinal bulb, and the total interval between stimulation and response  $\frac{1}{20}$  second. This is the shortest known reflex time on man, and it is noteworthy that stimulation of one eye causes simultaneous reaction of both eyelids, there being no time lost across the middle line (w.). We shall see that on the frog it may be shown that a crossed reflex suffers more delay in the cord than an uncrossed reflex. To determine the spinal delay on mammalia it would be necessary to measure the undoubtedly spinal reflex of animals after division of the cord, or that of a paraplegic patient. The latter observation has not been made; with regard to the former, Franc and Pitres give in the case of the dog  $\cdot 022$  to  $\cdot 040$  sec. for a reflex on the same side,  $\cdot 048$  to  $\cdot 058$  for a reflex on the opposite side.

*Adaptive or 'psychical' reaction.*—To what degree a brainless frog is superior to a brainless dog or man may best be realised by the following experiment on decapitated frogs. The frog is suspended, the side of its body is touched with strong acid; it raises the leg of the same side, and makes wiping movements at the spot of irritation; the leg of that side is held or removed, the animal may then curl the other leg round and wipe the spot of irritation. From these, among other observations, Pflüger was led to apply the term 'psychical' to the spinal function of the frog; and although the term may appear somewhat extravagant, it serves to call attention to the extremely complicated, and, to all appearance, discriminative and appropriately adapted reactions of the spinal centres in the lower vertebrate animals.

*Action of strychnine.*—Strychnine causes great exaggeration of the excitability of the spinal cord of all vertebrates. A man, a dog, a rabbit, or a frog, having received a poisonous dose of strychnine, dies in tetanus with the limbs extended and all the muscles of the body firmly contracted. In the case of a mammalian animal, the tetanus involves arrest of the movements of respiration and circulation, and death is definitive, the rigid state of the muscles passing off in a few moments, because the grey matter of the cord quickly ceases to act. In the case of a frog, even after a considerable dose, the animal remains stiff for hours, but is not in reality dead; cutaneous respiration is sufficient, circulation continues, and the spinal cord remains active. That the contracted condition of the muscles is entirely due to the spinal cord may be proved at once by destroying the cord, when the tetanus gives way and the animal becomes flaccid. That the exaggerated action does not in any degree depend upon increased excitability of either nerve or muscle may be ascertained by testing them with induction shocks; to this end it is advisable to protect one limb from the drug by ligature, and compare its nerve and muscle with those of the opposite, unprotected limb, when it will be found that, so far from being more excitable, the muscles of the unprotected limb are much less excitable than those of the protected limb; they exhibit, in fact, signs of having been fatigued by the exces-

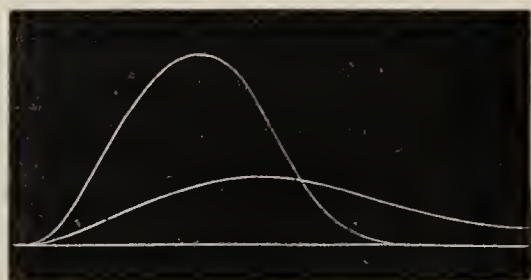


FIG. 273.—NORMAL CONTRACTION AND REDUCED CONTRACTION BEFORE AND AFTER STRYCHNINE TETANUS. DIRECT EXCITATION.



sive action into which they have been put by the spinal cord (fig. 273) (w.).

Another noteworthy feature in the strychnine tetanus of a frog is, that the contraction is not (at any rate at the outset of intoxication) continuous and unbroken, but serrated, *i.e.* the muscular tetanus is incomplete, with a contraction-frequency

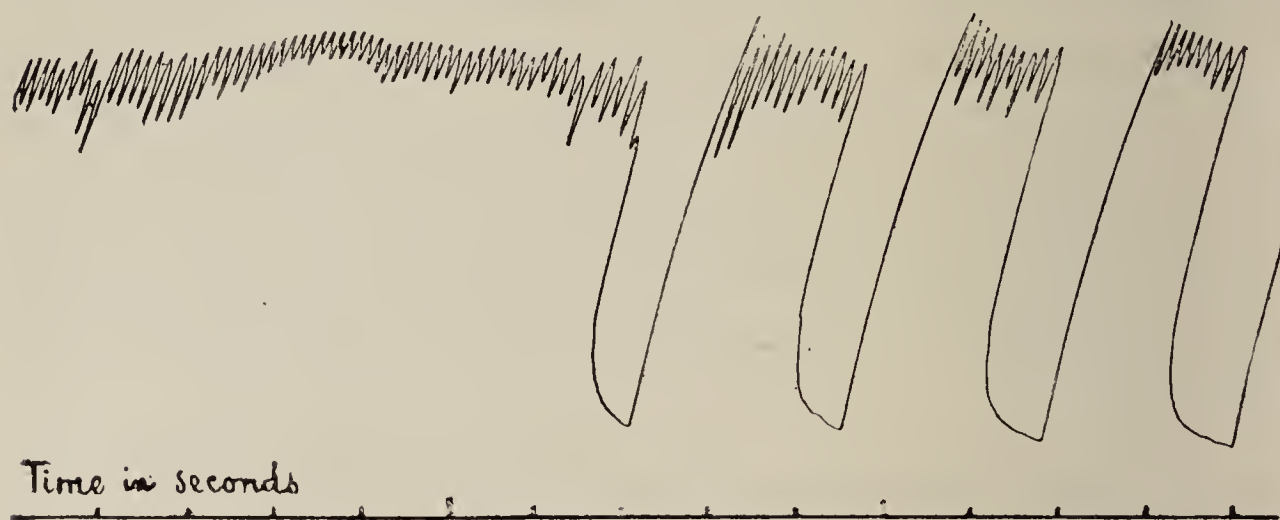


FIG. 274.—RECORD OF STRYCHNINE TETANUS; FROG.

of 8 to 10 per second—indicating that the spinal cord is discharging impulses at this rate.

Several further facts relating to the action of strychnine on the cord are studied on frogs. Summation is abolished, a single weak stimulus is as effectual as a series of stimuli; diffusion of stimulation is excessive, the excitation extends indiffe-

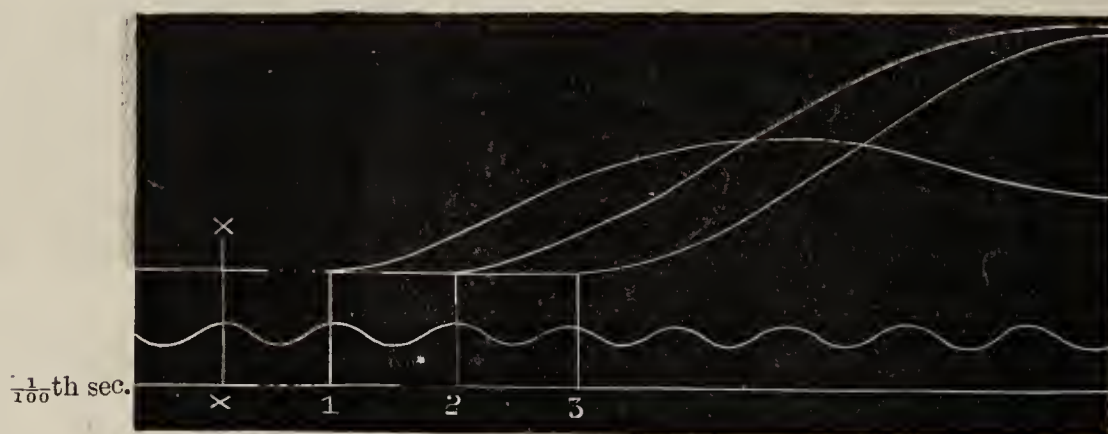


FIG. 275.—FROG.

Lost times of (1) a direct muscular contraction; (2) of a simple reflex contraction; (3) of a crossed reflex contraction; x marks the moment of excitation.

rently in all directions in the grey matter, and cannot be shown to extend transversely and longitudinally in proportion to its strength; in short, we may use for stimulation of the strychninised cord the same phrase as that by which we characterised the response of a quiescent heart (p.91)—‘all or nothing’; the stimulation produces no effect or a complete one, consisting in a convulsion of all the muscles of the body.

It might be expected that with an exaggeration of excitability such as that just described we should find the reflex time shortened. But the opposite is the case; although excitability is increased, the reflex time is prolonged, and this to such a degree that we can analyse the phenomenon on strychninised frogs far more minutely than is possible in a normal condition.

If on a slightly strychninised frog we measured the lost times—(1) of direct excitation, (2) of reflex reaction of the excited limb, (3) of reflex reaction of the opposite limb, (4) of reflex reaction of the limb above, we should find that (2) is about  $\frac{1}{100}$  second longer than (1), that (3) and (4) are respectively about  $\frac{1}{100}$  second longer than (2)—*i.e.* that the time lost in transverse and in longitudinal diffusion in the cord is about the same as that of a simple reflex process (w.). In a more deeply intoxicated frog, notwithstanding the greater excitability, the lost time is much increased, that of the simple reflex sooner and to a much greater extent than that of the transverse or longitudinal diffusion; the simple reflex time may be increased ten-fold, while the diffusion time is only doubled.

**Centres of the spinal cord and bulb.**—The actions ordinarily administered by the spinal cord are of various characters, and the spinal cord is credited with numerous centres ‘for’ various actions. In passing these in review it will be convenient at the same time to consider the central functions subserved by the spinal bulb, which likewise administers a variety of actions of a reflex or of an automatic character. The cord and bulb are in this connection to be regarded as of co-ordinate rank, constituting conjointly a medullary axis of grey matter—medulla oblongata and medulla spinalis—which acts as the intermediate between the brain and the periphery.

The spinal cord is credited with the following centres:

Movements of limbs . . . . .	page 503
Musculo-tonic . . . . .	„ 504
Respiratory . . . . .	„ 148
Cardiac accelerator . . . . .	„ 107, 111
Vasomotor . . . . .	„ 111
Sudorific . . . . .	„ 246
Cilio-spinal . . . . .	„ 448
Genito-spinal . . . . .	„ 110
Ano-spinal . . . . .	„ 170
Vesico-spinal . . . . .	„ 240
Trophic . . . . .	„ 359



The spinal bulb is credited with :

Respiratory	{ expiratory inspiratory }	. . . . .	page 148, 507
Vasomotor	{ constrictor dilatator }	. . . . .	„ 100, 110
Cardiac	{ accelerator inhibitory }	. . . . .	„ 100, 110
Centres of phonation and articulation ; coughing and			
sneezing	. . . . .	. . . . .	„ 159
Sucking	. . . . .	. . . . .	„ —
Mastication	. . . . .	. . . . .	„ 164
Deglutition	. . . . .	. . . . .	„ 166
Vomiting	. . . . .	. . . . .	„ 170
Co-ordinating	. . . . .	. . . . .	„ 503
Convulsor	. . . . .	. . . . .	„ 503
Closure of eyes	. . . . .	. . . . .	„ 498
Dilatation of pupil	. . . . .	. . . . .	„ 449
Salivary	. . . . .	. . . . .	„ 182
Sudorific	. . . . .	. . . . .	„ 246
Diabetic	. . . . .	. . . . .	„ 244

The expression centre '*for*' such and such an action is to be deprecated ; it implies a mapping out of the bulb and cord into departments far more definite and artificial than the reality, and an illegitimate dictation of final causes. And this artificial scheme is pushed to the verge of absurdity when centres such as the 'convulsive' or 'vomiting' centre are invented, centres for sneezing, laughing, crying, sucking, coughing, &c. Such 'centres' as these do not deserve the name. Certain actions are carried on by groups of associated muscles, served by nerves, which spring from different parts of the bulb and cord ; these different parts are anatomically centres, but it is an abuse of language to call them centres *for* such actions. We are not justified in speaking of a centre for muscular tone, and of a centre for movements of the limbs, as if they were separate real organs ; cilio-spinal, genito-spinal, ano-spinal, vesico-spinal, convey a notion of definiteness far in excess of actual facts, which are simply to the effect that nerves to the iris, genitals, rectum, and bladder are connected with the spinal cord, and that certain reflex actions of these various parts require the integrity of the cord, and of the nerves connecting them with it. So, again, with regard to the so-called sudorific and trophic centres, and the somewhat less indefinite cardiac, respiratory, and vasomotor centres in the bulb and cord ; these are one and all indefinite organs, and not to be regarded in the light of

anatomically definable nuclei of grey matter. Nerves, that are channels of sudomotor, vasomotor, and respiratory acts are connected with certain segments of the medullary axis; this is the simple anatomical fact, to a knowledge of which it is no addition to say that they are the specific centres for certain functions. With such or some such reservations made, we may consider in detail the different kinds of actions that are controlled from the medullary axis.

*Movements in general.*—Simple reflex movements of the limbs may result from peripheral stimuli; complex, orderly, co-ordinated movements, such as those of locomotion, may also be performed automatically. Movements of both these kinds are governed by the spinal cord, and are presumably administered by it; it is chiefly on the lower animals, however, that the experimental proof has been given of the performance of co-ordinated movements after the removal of the brain and bulb. Various notable experiments have been made on frogs and on snakes in this connection. A decapitated frog, possessing no nervous centre above the spinal cord, is a living machine that moves only when it is set in motion by a stimulus—when it does move, however, it exhibits indications of a high degree of complexity. The headless frog makes definite and appropriate efforts to remove an irritant applied to its skin; if it fail with one leg, it makes attempts with the other. A decapitated snake likewise makes highly complex movements: twines round a stick, bends its body away from a hot poker. On the higher animals even highly complex movements may still be performed: ducks and ostriches have been known to run about after decapitation. Mammalia, however, exhibit little beyond simple reflex movements, though even in their case the movements are such as to suggest self-defence as their *raison d'être*.

There is no reason whatever for admitting the existence of any distinct *co-ordinating centre* in the bulb or cord. When the co-ordinate and measured movement of a group of muscles occurs, due order and measure of outgoing impulses are of primary necessity; a co-ordinate character is not imposed upon the action of some centres by other specially co-ordinating centres. There are no grounds for admitting co-ordination as a special function of special centres. Nor is there any reason for admitting the existence of any distinct *convulsion centre* in the medulla or cord. When a convulsive action occurs, out-



going impulses are without order and measure; an inco-ordinate and excessive character is not imposed upon the action of some centres by any other specially convulsive centre. There are no grounds for admitting convulsion as the special function of any special centre.

The *musculo-tonic* action exercised by the spinal cord keeps the whole muscular system in a state of slight contraction or tone. There is no reason for attributing such action to special centres; it is in any case not to be regarded as anything beyond a slight and continuous motor discharge, and there is no reason for regarding it as different in kind or origin from the stronger and discontinuous motor discharges that cause muscular contractions. The very existence of a musculo-tonic action has been questioned, but is, nevertheless, to be regarded as established by sufficiently conclusive experiments. Brondgeest's experiment to demonstrate muscular tonus consisted in the simple suspension of a frog after section of one sciatic nerve; the normal limb remained flexed, the other limb hung loose. Cyon performed the experiment after section of the posterior roots of the spinal nerves that go to form the sciatic, with a similar result, and has shown that the tonic flexion ceases when the limb is deprived of its skin; on the other hand, Sherrington found that the tone of muscles was increased by stimulating afferent nerves issuing from antagonist muscles. These are proofs that muscular tone is a continuous reflex action governed by centripetal influence.

The phenomenon of so-called *tendon-reflex* on man bears a close relation to muscular tonus. It consists in the sudden con-

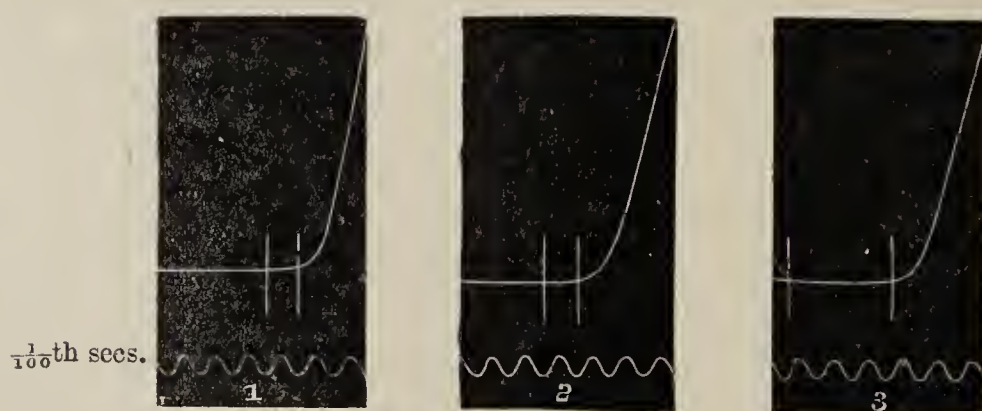


FIG. 276.

Time measurements of (1) a direct contraction, (2) of the tendon phenomenon, (3) of a reflex contraction (on the rabbit).

traction of a muscle when its tendon is smartly struck, and is best known in the case of the rectus femoris when the liga-

mentum patellæ is struck. It is not a reflex contraction, for its latent period is about the same as that of muscular contraction caused by direct electrical stimulation, *i.e.* much too short to be acceptable as the time of a reflex contraction (w.); but some influence upon muscle of a reflex nature is its indispensable condition, for if the nerves supplying the muscle be cut, or if only their motor roots be cut, or if only their sensory roots be cut, the reaction is abolished; *i.e.* the integrity of the reflex arc is a necessary condition of the reaction. The contribution of central action is further signified by the fact that the phenomenon is intensified during voluntary action, depressed during sleep or anæsthesia, and in anæmia of the cord; and Sherrington finds that it is abolished by centripetal excitation of nerves coming from the particular muscles (or from their connective tissue) that are of antagonistic action. It has great value as a clinical test, especially in the diagnosis of locomotor ataxy; its abolition is an early and very characteristic sign of the onset of this disease. The opposite change, *viz.* an exaggeration of the reaction, usually occurs as a sign of descending degeneration of the cord in consequence of lesions of the cord or brain, and the exaggeration is generally accompanied, or very shortly followed, by a state of permanently increased muscular tonus—the ‘*contracture*’ of clinical medicine. ‘Contracture,’ if not of very old standing, is cut short by section of the motor nerves, and disappears during sleep or anæsthesia, and at death. These points show that it is of central origin; it may occur in an incomplete form as a more or less prolonged ‘stiffening’ brought on by voluntary effort.

The tonic contraction of the sphincters of the rectum and bladder is probably of a similar nature, *viz.* a reflex spinal tonus; incontinence of fæces and of urine are among the symptoms of disease of the lumbar portion of the cord, and Heidenhain has shown experimentally on rabbits that the resistance of the vesical sphincter to intravesical pressure is greater during life than after death, thus proving that the closure is not solely due to elasticity, but to elasticity reinforced by tonic contraction.

*Equilibration.*—Among the numerous natural groups of movements that enter into the motor conduct of the body, those of equilibration and of locomotion especially attract attention by their universal importance and constant necessity. They are reflex adjustments to peripheral sensations—visual, labyrinthine,



cutaneous, and muscular ; and there is a disposition to regard these adjustments as being reflex in more *detailed* manner than other groups of co-ordinate movements, such as those of speech, or of writing, or of special crafts. It is considered that the state of tone and of contraction is, item by item, in direct reflex response to peripheral muscular tension, exciting a muscular sense by the channel of afferent nerves from the muscles themselves, or from their tendons, or from the articulating surfaces. This may be so, but it is not proved ; it is not even probable. No doubt reflex adjustments are continually ebbing and flowing in the maintenance or alterations of muscular attitude, and the existence of a reflex 'tonus,' if not conclusively proved, is highly probable. But the assumption that in a series of movements each individual movement is a reflex spinal response to a pull of the muscle, or of the tendon, is inadmissible ; in the large, slow movements of locomotion, and still more in the small, rapid movements of vocalisation, we must suppose that the constituent items in the co-ordinate series are measured out in due proportion and sequence in the emissive organ, independently of discrete peripheral calls through 'muscular sense.' There are also reasons for admitting that a direct peripheral response of muscle to extensile stimuli contributes to that harmonised action of antagonistic muscles which insures their smooth and steady service (*vide* p. 344). Bearing these points in mind, we are free to recognise the great importance of central guidance by peripheral conditions. A normal man walking perfectly, with his eyes open, with a normal state of his semicircular canals and of his plantar touch-organ, and with a normal sensory innervation of his tendons, ligaments, and articular surfaces, will walk less or more imperfectly if one or more of these sources of information is perturbed or stopped. But this is not all, and in the co-ordinate function of any synergic group of muscles at least three conditions must be fulfilled :—(1) Impulses must be emitted from the centre in appropriate manner and degree ; (2) the muscles must be 'in touch' with the spinal axis, duly responsive to the reflex ebb and flow arising from altering peripheral states ; (3) they must be in touch with each other, directly reacting against that passive extension which antagonist muscles undergo from each other. If the first condition be at fault, the result is a disorder of movement, of which 'stammering' is the type ; if the second and third are out of measure (and they are so closely associated as usually to go right or wrong together), we witness 'ataxic'

effects, of which locomotor ataxy is the most familiar and extreme example.

The nerves from the vestibule and semicircular canals form the vestibular nerve, which is anatomically traceable by the ventral root of the eighth nerve to the lateral portion of the auditory nucleus in the spinal bulb. From this station onwards the destination of the nerve is not so clear; tracts from the nucleus are said to pass to the cerebellum by way of the restiform body. The principal end-organ of the vestibular nerve is formed by the bristle-cells of the so-called 'acoustic' crests of the ampullæ. The semicircular canals are probably not auditory in function, but 'equilibratory'—*i.e.* they are the origin of sensations caused by pressure-effects of the endolymph in the ampullæ, and

contributing to the maintenance of normal muscular equilibrium. The directions of the three canals are very nearly in those of the three dimensions of space, and the planes occupied by the six canals are approximately as indicated in the diagram. Thus a rotation of the head clock-wise would give rise to greater pressure of

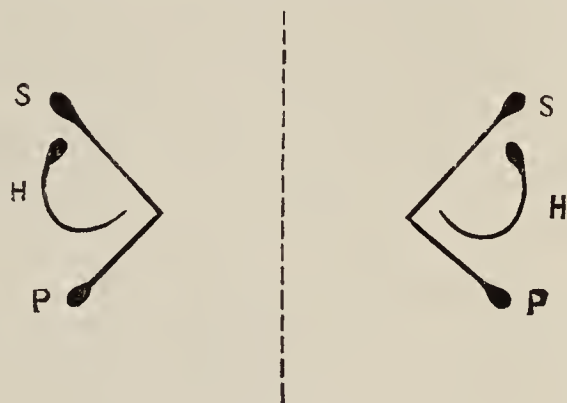


FIG. 277.

Diagrammatic horizontal section through the head to illustrate the planes occupied by the semicircular canals.

endolymph in the ampulla of the right horizontal canal; a sudden cessation or reversal of rotation would give greater pressure in the ampulla of the left horizontal canal (Crum Brown). Clinical records are appealed to in support of the view that the integrity of the semicircular canal is essential to equilibration. 'Menière's disease' has as its prominent features deafness, giddiness, and staggering, presumably due to irritative lesion of the internal ear. And among the experimental causes of forced movements, excitation of the semicircular canals plays a prominent part. On the other hand, it should be mentioned that most careful experiments made on cartilaginous fishes have failed to demonstrate any connection between the semicircular canals and the co-ordination of movements (Sewall, Steiner).

*Respiratory centre.*—The movements of respiration constitute a very definite act, carried out by muscles that receive their nerves from a certain definite part of the bulbar centre, which is influenced through afferent nerves, and by the state of the blood.



The term 'respiratory centre' has thus a far better justification than, for instance, such terms as 'convulsive' centre, or 'vomiting' centre. It is anatomically that part of the medullary axis from which originate the phrenic and pneumogastric nerves, and is situated at the lower part of the bulb and upper part of the cord.

Definite nuclei of grey matter have, however, never been experimentally isolated as a respiratory centre or centres. Destruction of the lower part of the spinal bulb by means of an instrument introduced between the occiput and the atlas causes sudden death by cessation of respiratory movement; and Flourens applied to this part the sensational name of 'nœud vital.' 'Nœud vital' and 'respiratory centre' have now for us an identical meaning, and signify a part of the bulb with which are connected nerves whose integrity is necessary to the continuance of respiratory movements, that is to say, to the continuance of life. The respiratory centre is situated in chief part in the bulb, in lesser part in the upper part of the spinal cord. Its mode of action may be characterised as reflex and automatic: reflex inasmuch as it may be temporarily increased or diminished by occasional centripetal impulses; automatic inasmuch as it appears to go on of itself, being in reality kept going by continuous stimuli. These continuous stimuli reach the respiratory centre in two ways—(1) the state of the blood—reduced blood is excitant, oxygenated blood is depressent of its excitability; (2) the pneumogastric nerves. But these matters have already been considered under the heading of Respiration. As regards the vasomotor actions of the spinal cord and bulb, we have already learned the main physiological facts in a previous section.

We may, indeed, bring our review of the specific central actions of the bulb and cord to a close; its purpose will have been sufficiently answered if the lists given on pp. 501–2—to which are appended the references to each particular subject—have served to bring into prominence the number and importance of the functions to which the integrity of the medullary axis is necessary. It should also be realised that the closely congregated nuclei of grey matter (see figs. 279, 280) located in its upper or bulbar expansion, are concerned in the performance of those fundamental functions which are of most immediate importance to the living body—respiration, deglutition, regulation of the heart and of the vessels. And we may in this connection conclude with a reference to the group of cardinal symptoms that

are characteristic of bulbar paralysis, *i.e.* of an impaired medulla oblongata; these are a progressive imperfection of articulation, of mastication, of deglutition, and of phonation—‘glosso-labio-laryngeal-pharyngeal’ palsy.

### THE CRANIAL OR BULBAR NERVES

It will be convenient at this stage to review the physiological anatomy and chief functions of the cranial, or so-called cerebral,

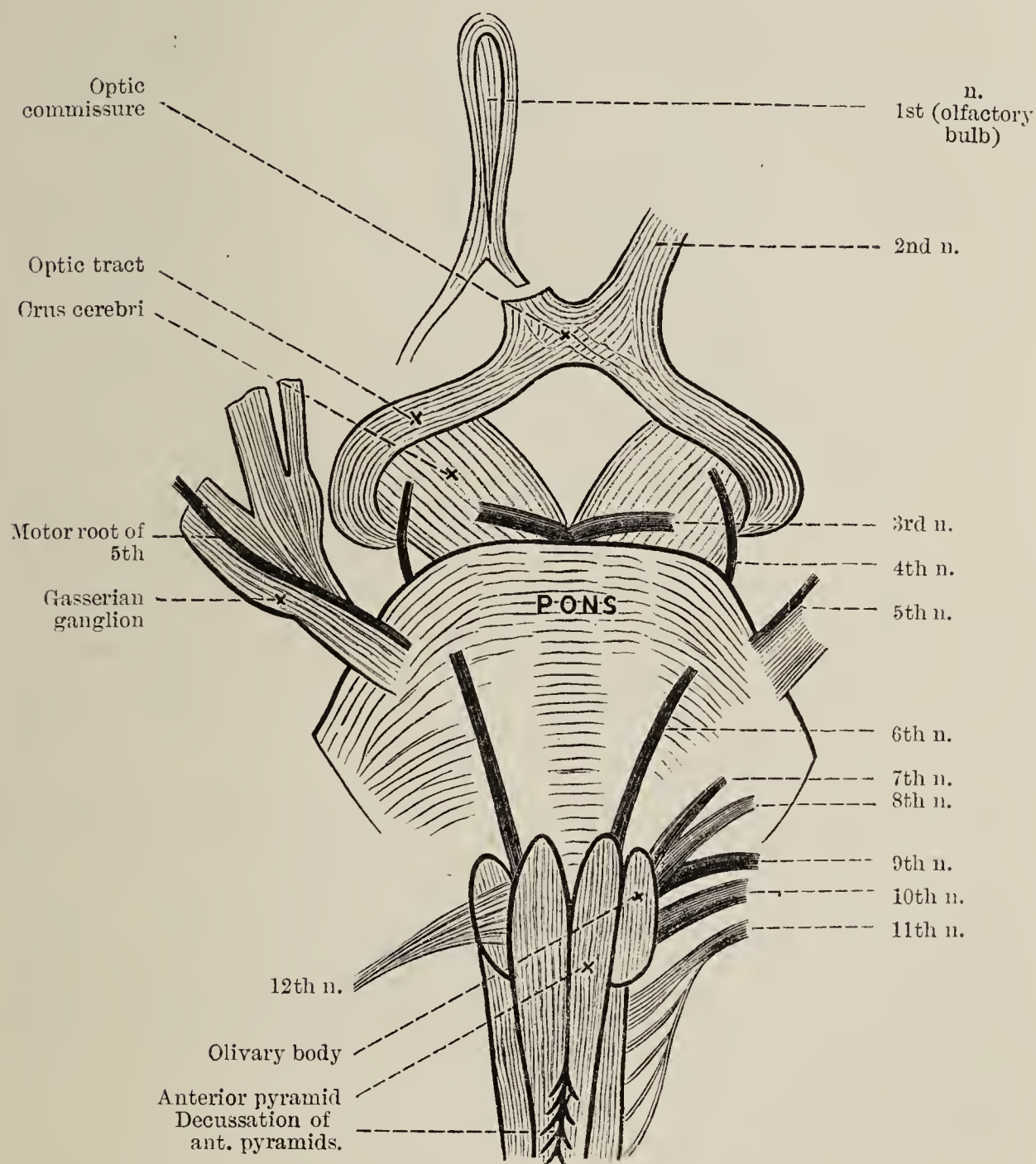


FIG. 278.—ANTERO-INFERIOR VIEW OF THE CRURA, PONS, AND BULB (Diagrammatic), TO ILLUSTRATE THE SUPERFICIAL ORIGIN OF THE CRANIAL NERVES.

nerves. These nerves, twelve on each side, emerge from the base of the brain at points indicated in fig. 278. From this, their superficial origin, their fibres are traceable to nuclei of grey matter, which are clustered together in the neighbourhood of the



fourth ventricle, as indicated in figs. 279 and 280. The cranial nerves are functionally connected with the cortex of the opposite side, and, as in the case of the spinal nerves, there are no direct fibres between cortex and the periphery; every fibre has at least one spinal cell in its course. Thus, in the case of cranial, as in that of spinal nerves, the path between cortex and periphery is

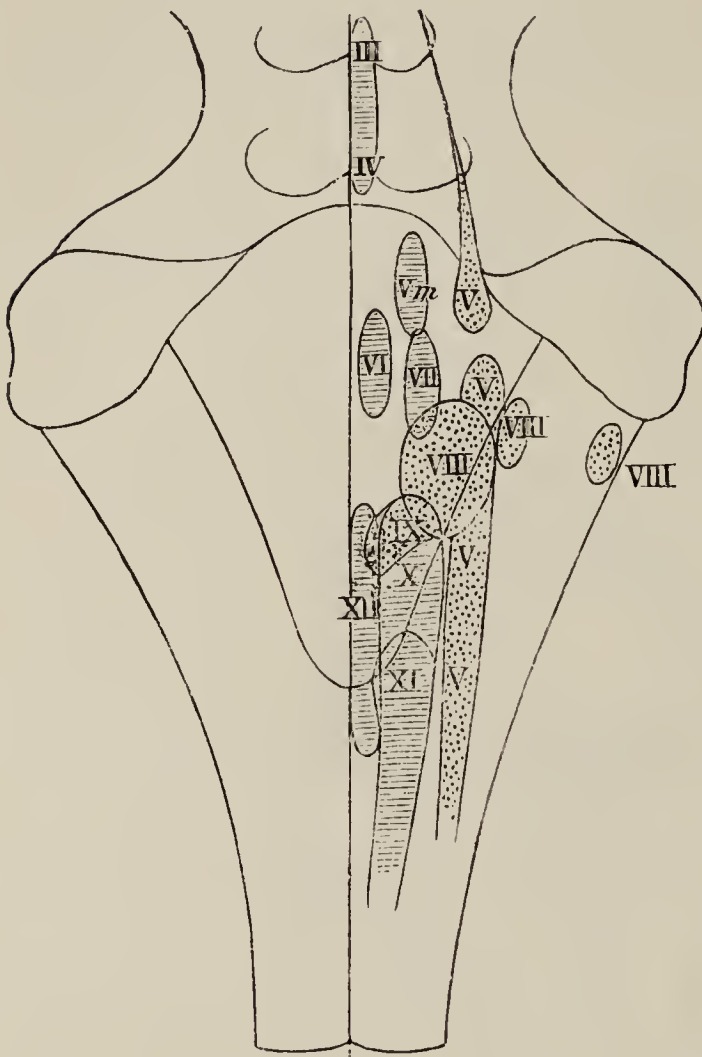


FIG. 279.—DIAGRAMS TO ILLUSTRATE THE POSITION OF THE BULBAR NUCLEI OF THE CRANIAL NERVES.

Posterior aspect or 'floor' of the fourth ventricle exposed by removal of the pons and cerebellum, and imagined as transparent. Motor nuclei indicated by horizontal lines, sensory nuclei by dots. Median group of motor nuclei, III, IV, VI, XII. Lateral group of motor nuclei, Vm, VII, X, XI. Sensory nuclei, Vs, VIII, IX.

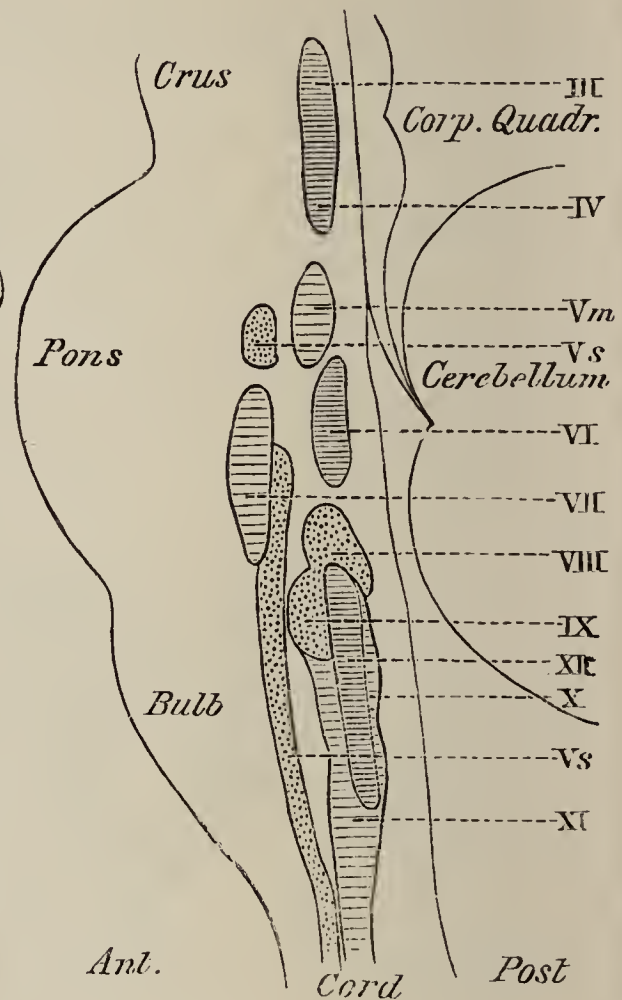


FIG. 280.—LATERAL VIEW OF THE RIGHT HALF OF THE BULB AND PONS EXPOSED BY A VERTICAL SECTION, AND IMAGINED AS TRANSPARENT. (After Erb.)

In this view the lateral group of motor nuclei, Vm, VII, X, XI, lie farther from the surface of section, and are indicated by lighter lines than the median group of motor nuclei, III, IV, VI, XII.

divisible into at least two stages; one between cortical and bulbar cell, and one between bulbar cell and end-organ (muscle or epithelium). Thus, properly speaking, all cerebral and spinal motor and sensory nerves (with the exception of the first and second) are cerebro-spinal as regards their course, the 'cerebral' nerves having their spinal relay in the bulb, the 'spinal' nerves

having theirs in the cord, in either case across the middle line; and if a classification is to be made into groups, that which most commends itself is into *spinal* and *bulbar* groups, the latter comprising the first twelve nerves, exclusive of the first and second, which occupy an exceptional position, but inclusive of the third and fourth, which, although arising from nuclei imbedded in the mid-brain, are in line, anatomically as well as functionally, with the sixth and twelfth nerves.

The cranial nerves are as follows :

Name.	Distribution.
1. Olfactory nerve .	From nose.
2. Optic „ .	From eye.
3. Oculo-motor nerve	To eye muscles.
4. Trochlearis . .	To superior oblique.
5. Trigeminal . .	From face, mouth, nose. To muscles of jaw.
6. Abducens . .	To external rectus.
7. Facial . .	To facial muscles.
8. Auditory . .	From ear.
9. Glosso-pharyngeal .	To pharynx. From tongue.
10. Pneumogastric .	To and from heart, lungs, viscera.
11. Spinal accessory .	To heart. To trapezius and sterno-mastoid. To larynx, pharynx, and œsophagus.
12. Hypoglossal . .	To tongue.

Traced back from their superficial to their deep origin, most of the cranial nerves are found to spring from nuclei of grey matter closely congregated in the spinal bulb and pons beneath the floor of the fourth ventricle. The second, or *optic*, nerves by their optic tracts have extensive connections with the basal ganglia; fasciculi of the optic tract are traceable from three sources of origin—the lateral corpus geniculatum, the anterior corpus quadrigeminum, and the optic thalamus (posterior tubercle or pulvinar). The third, or *motor oculi*, and the fourth, or *trochlearis*, have their nuclei of origin in the corpora quadrigemina, close to the Sylvian aqueduct. The fifth, or *trigeminal*, has three distinct nuclei of origin; the median and smallest of these, in the floor of the fourth ventricle, is the ‘motor nucleus,’ the other two are the ‘sensory nuclei,’ and give origin to the ‘ascending’ and ‘descending’ roots of the nerve. The sixth, or *abducens*, has its nucleus of origin in the floor of the fourth ventricle. The nucleus of the seventh, or *facial*, lies deeper, below the floor of the fourth ventricle, in the substance of the pons. The three *auditory* nuclei (eighth nerve)—median, lateral, and accessory portions, occupy a lateral situation in the upper



part of the floor of the fourth ventricle; the more superficial or median, or principal nucleus, and the smaller accessory part (which bears some resemblance to a spinal ganglion), give rise to the cochlear division; the lateral or deeper, or superior portion, gives rise to the vestibular nerve. The nerve of Wrisberg (pars intermedia of the seventh), from which the chorda tympani is formed, is also said to arise from the accessory nucleus. The ninth, tenth, and eleventh nerves—*glosso-pharyngeal, vagus, and accessory*—arise from an elongated nucleus of grey matter laterally situated in the lower part of the medulla and upper part of the spinal cord. The twelfth, or *hypoglossal* nerve, has its nucleus of origin superficially in the lower part of the bulb, close to the middle line.

The *first, or so-called olfactory nerve*, is in reality a lobe of the brain; the true olfactory nerves are the eight or ten filaments that are connected with it. This lobe is the central organ of smell, the filaments are afferent of impulses caused by stimuli affecting the olfactory epithelium contained in the nasal mucous membrane.

The *second, or optic nerve*, is the continuation of the optic tract, which is in connection with the corpora quadrigemina. The peripheral organ is a particular layer of the retina (rods and cones), and its cortical terminus is the occipital region. Section of the optic nerve causes blindness of the corresponding eye, excitation of its central end causes the sensation of light (subjective). It is an afferent nerve, conveying the impulses which in consciousness give rise to the sensation of vision.

The fibres of the optic nerve cross the middle line at the optic chiasma; in the lower animals and in many mammalia (rat, guinea-pig, sheep, pig, horse, goat, deer) the decussation at the chiasma is complete; in other mammalia (rabbit, dog, cat, monkey, man) it is partial, each optic nerve giving a large proportion of fibres to the optic tract of the opposite side, and a small proportion of fibres to the optic tract of the same side; section or lesion of one optic tract in the case of man and of the higher mammalia causes paralysis of both retinæ (hemianopia) on the side of the lesion—‘bilateral homonymous hemianopia,’ p. (543).

The *third, or motor oculi*, supplies all the muscles of the eyeball (with the exception of the superior oblique and external rectus), the elevator muscle of the upper eyelid, the iris, and the ciliary muscle. Section of the third nerve paralyses these, the

eyeball is protruded and squints downwards and outwards, the upper eyelid drops, the pupil is enlarged, the eye cannot be accommodated. Excitation of its peripheral end reverses all these effects: the eyeball squints upwards and inwards, the upper eyelid is raised, the pupil contracts, and there is spasm of accommodation. Both the above-mentioned groups of signs are of common clinical occurrence in consequence of paralysis or of irritation of the third nerve.

The *fourth, or trochlearis*, supplies the superior oblique muscle of the eyeball. Its section or paralysis causes the eye to squint inwards and upwards.

The *fifth, or trigeminal*, is a mixed nerve composed of a larger sensory portion and of a smaller motor portion; the former is a sensory nerve of the face, of the conjunctiva, of the teeth, of the mucous membrane of the nose, mouth and tongue; the latter is the motor nerve of the muscles of mastication, of the tensor veli palati, and of the tensor tympani muscles. Nerve-fibres having direct control over nutrition (trophic) are assumed to be present in the fifth nerve, but the existence of these trophic fibres has not been conclusively determined (p. 361). The chief consequences of section of the fifth are:—(1) Paralysis of common sensation in its area of distribution, accompanied with impairment of smell and of taste; (2) paralysis of the muscles of mastication; (3) disorders of nutrition, of which the most notable effect is inflammation of the cornea, leading to ulceration, and ultimately to complete destruction of the eyeball. The branches of the fifth contain numerous vasomotor fibres derived from the sympathetic. These are mainly vaso-dilatator in kind.

The *sixth, or abducens*, supplies the external rectus muscle of the eyeball. Its section or paralysis causes the eye to squint inwards.

The *seventh, or facial*, is the motor nerve of the muscles of the face, *i.e.* it is the nerve that governs facial expression. By virtue of its chorda tympani branch it is also the channel of secreto-motor and of vaso-dilatator impulses; these are, however, attributable to sympathetic fibres. The gustatory fibres of the chorda tympani are derived from the glosso-pharyngeal. If the seventh nerve of one side is paralysed (Bell's palsy), the face on that side is smooth and expressionless, the eyes cannot be closed, lip movements, such as whistling or the articulation of labial consonants, are interfered with, and masticated food tends to collect between the cheek and gum, owing to paralysis of the



buccinator; in consequence of the unopposed action of the muscles of the opposite or sound side, the face is drawn towards that side, particularly when the patient is made to smile or laugh. If a lesion of the facial occurs within the Fallopian canal, so as to involve the chorda tympani, taste and salivary secretion will be affected. Section of the facial in young animals interferes with the nutrition of the muscles and bones of that side.

The *eighth, or auditory*, is the nerve of hearing, and plays an important part in equilibration. The peripheral organ is the internal ear, the auditory portion proper being in connection with the cochlea, the equilibratory portion in connection with the semicircular canals. Section of the auditory nerve causes deafness, giddiness, and staggering gait. Excitation of its central end gives rise to the subjective sensation of sound. *Menière's disease*, of which the characteristic symptoms are giddiness and staggering gait, has been found to be associated with affections of the semicircular canals.

The *ninth, or glosso-pharyngeal*, is a mixed nerve; it is the sensory and gustatory nerve of the posterior third of the tongue and adjoining mucous membrane of the mouth and pharynx, and the motor nerve of the stylo-pharyngeus and of the middle constrictor of the pharynx. Excitation of the central end of the glosso-pharyngeal inhibits the act of deglutition, and may excite vomiting.

The *tenth, vagus, or pneumogastric*, is a mixed nerve, with which we must include the internal branch of the spinal accessory, or eleventh nerve. The pharyngeal branches forming the pharyngeal plexus are the motor nerves of the levator palati and of the three constrictors of the pharynx. The *superior laryngeal* branch is the sensory nerve of the mucous membrane of the larynx, and the motor nerve of one of the laryngeal muscles—the cricothyroid. The *depressor nerve* from the heart may have its course in the vagus trunk or in its superior laryngeal branch. After section of both superior laryngeal nerves the mucous membrane of the larynx is insensible, food or saliva can pass through the larynx without exciting coughing, and subsequently cause inflammation of the lungs. Stimulation of the central end of the superior laryngeal excites coughing, arrests respiration, and excites deglutition; it usually causes a reflex rise of blood-pressure, but may (if it includes the depressor) cause a reverse effect. The *inferior laryngeal* branch is the chief

motor nerve of the muscles of the larynx. The *cardiac branches* contain the inhibitory fibres to the heart, and the depressor fibres from the heart; they also contain accelerator fibres derived from the sympathetic. The *pulmonary branches*, forming the pulmonary plexus, contain sensory fibres from the bronchial mucous membrane, and motor fibres to bronchial muscle. They also contain fibres the excitation of which causes modification of respiratory movements. The *œsophageal branches*, forming the œsophageal plexus, the *gastric branches*, forming the gastric plexus, and the *intestinal branches*, are motor and sensory.

As regards the functions of the two portions of the vagus, viz. vagus proper and accessory, the former is mainly an afferent nerve, formed by fibres from the gastro-intestinal tract, from the heart, lungs, and larynx; the latter is exclusively an efferent nerve, giving off motor fibres to the levator palati, larynx and œsophagus, inhibitory fibres to the heart, and in all probability motor fibres to the bronchial muscle, to the stomach, and to the intestines.

*Experimentally*, section of the vagus high up in the neck causes acceleration of the heart-beat, impediment to deglutition, paralysis of laryngeal muscles, and—at a later date—inflammation of the lungs. Excitation of the peripheral cut end causes arrest of the heart's action, contraction of the larynx and of the œsophagus, and—but not so evidently—contraction of the bronchi, of the stomach, and of the intestine. Excitation of the central end causes reflex acceleration of respiratory movements, reflex inhibition of the heart, and reflex fall of blood-pressure.

The *eleventh, or spinal accessory*, nerve divides into two branches—the internal branch, just considered, which accompanies the vagus proper, and the external branch, which is the motor nerve of the sterno-mastoid and trapezius muscles.

The *twelfth, or hypoglossal*, is the motor nerve of the lingual muscles, and of the genio-hyoid and thyro-hyoid muscles. Paralysis of the hypoglossal interferes with the movements of the tongue, which, when protruded, deviates towards the paralysed side.

The **sympathetic** system of nerves has already been frequently referred to; it is a main channel of vascular and visceral nerve-control, and from a physiological standpoint its anatomy has been sufficiently described under the heading of vasomotor nerves (p. 101). We allude to it in this place on account of its origin from the grey matter forming the central axis of the spinal bulb



and cord ; the question formerly debated, whether the sympathetic system is the independent companion of the cerebro-spinal system, or whether it is a dependent province of that system, may at the present day be regarded as finally answered in the sense of the second alternative. The clearest and most definite proof that sympathetic nerves are under spinal control is afforded—(1) by the discovery that irido-motor fibres take origin from the spinal cord and bulb, (2) that vasomotor fibres have a similar origin. Confirmatory evidence is derivable from the facts (3) that the fine medullated fibres, which there are reasons for considering to be vasomotor, are traceable from the spinal cord into sympathetic nerves ; and (4) that no reflex functions can be attributed to any sympathetic ganglion. As pointing in the same direction, although the fact itself is not uncontested, nor in any case demonstrative whether or not the sympathetic exercises independent function in the adult state, we may allude to its embryonic origin as an outgrowth from the medullary axis (p. 584). The old view, according to which the sympathetic system of vertebrata was regarded as homologous with the entire nervous system of invertebrata, is entirely abandoned ; the homology is between the entire nervous systems of the two sub-kingdoms respectively.

## CHAPTER XV

## THE BRAIN

## PAGE

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- 518 **The hemispheres:** Cortex cerebri—Corona radiata—Internal capsule—Crura cerebri—Corpus callosum—The pyramidal or motor tract—The basal ganglia—Corpus striatum—Optic thalamus—The cerebellum—Corpora geniculata—Corpora quadrigemina.
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- 522 Functions of basal ganglia; of the cerebellum—Forced movements.
- 527 The cortex of the brain.
- 528 Arteries of the brain—The Sylvian artery—The lenticulo-striate artery.
- 530 **The functions of the brain.**  
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- 534 **Cortical localisation of function.**  
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- 545 Epilepsy—Epileptogenous foci.
- 547 Aphasia—Motor aphasia—Sensory aphasia—Word-blindness—Word-deafness—Recovery from aphasia—Deaf-mutes.
- 552 **The personal equation or reaction time.**
- 554 **Ratio between stimulus and sensation—Weber-Fechner law.**
- 556 Weight-discrimination—The muscular sense.
- 558 **Sensory judgments—Illusions—Contrast theories—Helmholtz, Hering.**
- 569 **Fatigue—Sleep—Hypnosis.**

*General plan of its structure.*—Without entering upon any detailed description of the brain as given in anatomical text-books, we shall in this place consider and insist upon such points only as have immediate bearings upon the knowledge of its functions.

The cranium contains the two cerebral hemispheres and their basal ganglia (corpora striata, corpora quadrigemina, optic thalami, corpora geniculata), the cerebellum, and the medulla oblongata. From the point of view of physiological anatomy we regard the medulla oblongata (= spinal bulb) as being spinal rather than cerebral; and we consider by itself the cerebellum, which, whatever its function may be, is not known to possess



any action that it is customary to call 'cerebral.' As regards the basal ganglia, our attention becomes directed to them, less on account of any definite functions that have been assigned to them, than because they occupy the isthmus of the brain, and are thus on or close to the path between brain and body. Our physiological study of the brain is thus simplified, and comes to be directed almost exclusively to the surface of the hemispheres, and to the tracts that connect them with each other and with

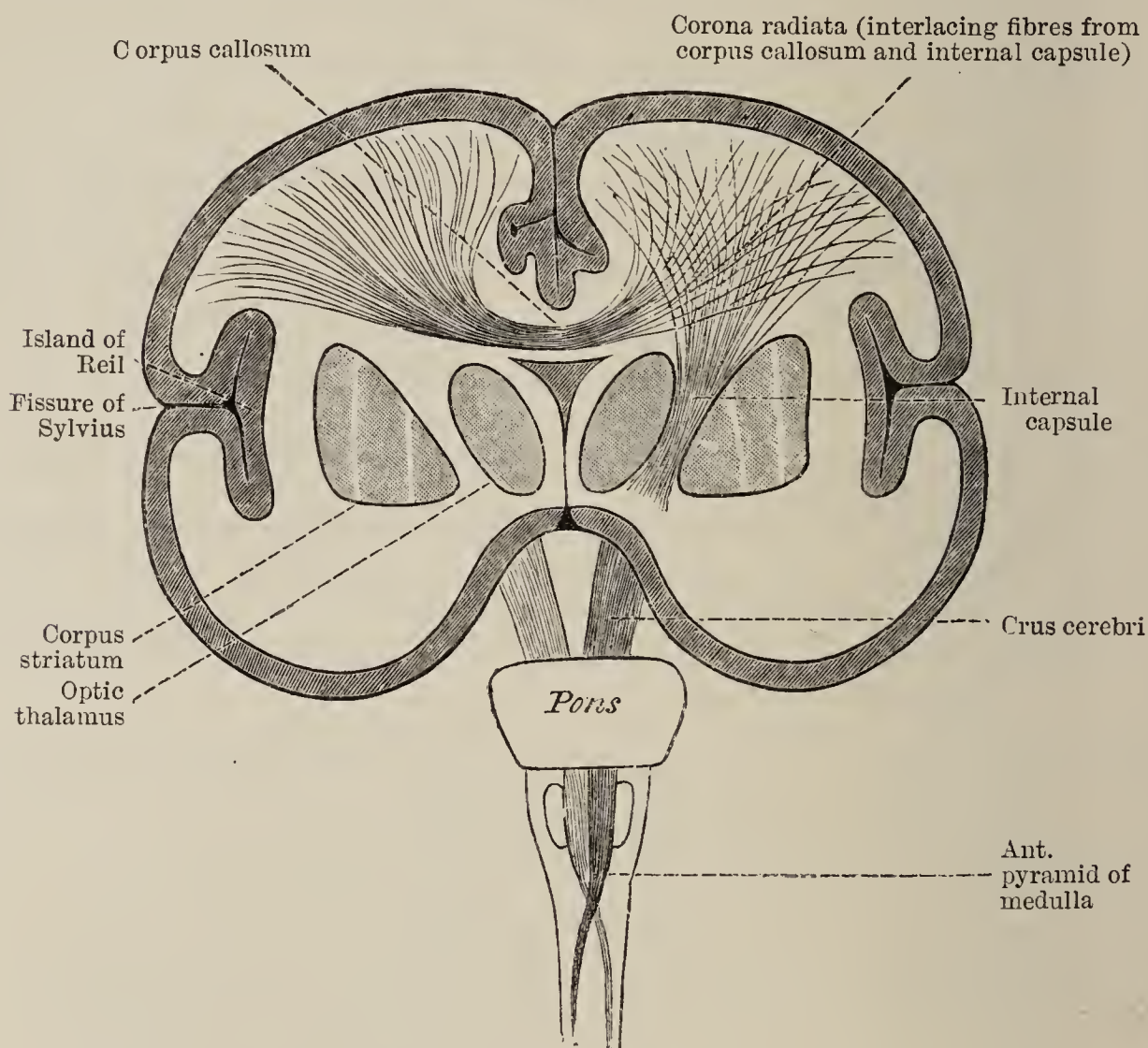


FIG. 281.—IDEAL VERTICAL SECTION THROUGH THE CEREBRO-SPINAL AXIS.

To illustrate the course of the pyramidal tract and its interlacement in the corona radiata with the commissural fibres of the corpus callosum.

the nerve-channels emerging from the base of the brain; these are the *crura cerebri* and the *cranial nerves*.

Viewed broadly, the brain is a mass of white matter, with nuclei of grey matter deeply imbedded in it, and with a sheet of grey matter, about one-fifth of a square meter in area and between 2 and 3 mm. thick, covering the folds, fissures, and convolutions of its surface. This superficial grey matter, or *cortex*, is the brain proper, the organ of sensation, judgment, and will; the white matter beneath leads into it sensory impulses, leads off from it

motor impulses, and is simply a conducting mass composed of bundles of medullated nerve-fibres, disposed in a fanlike manner between the isthmus of the brain and the grey matter of the cortex, while below the isthmus it is continued as the white columns of the bulb and cord. This mass of white matter is known as the *corona radiata*; at the base of the brain, between the corpus striatum and optic thalamus, it forms a conical sheet of white matter known as the *internal capsule*, and proved to constitute

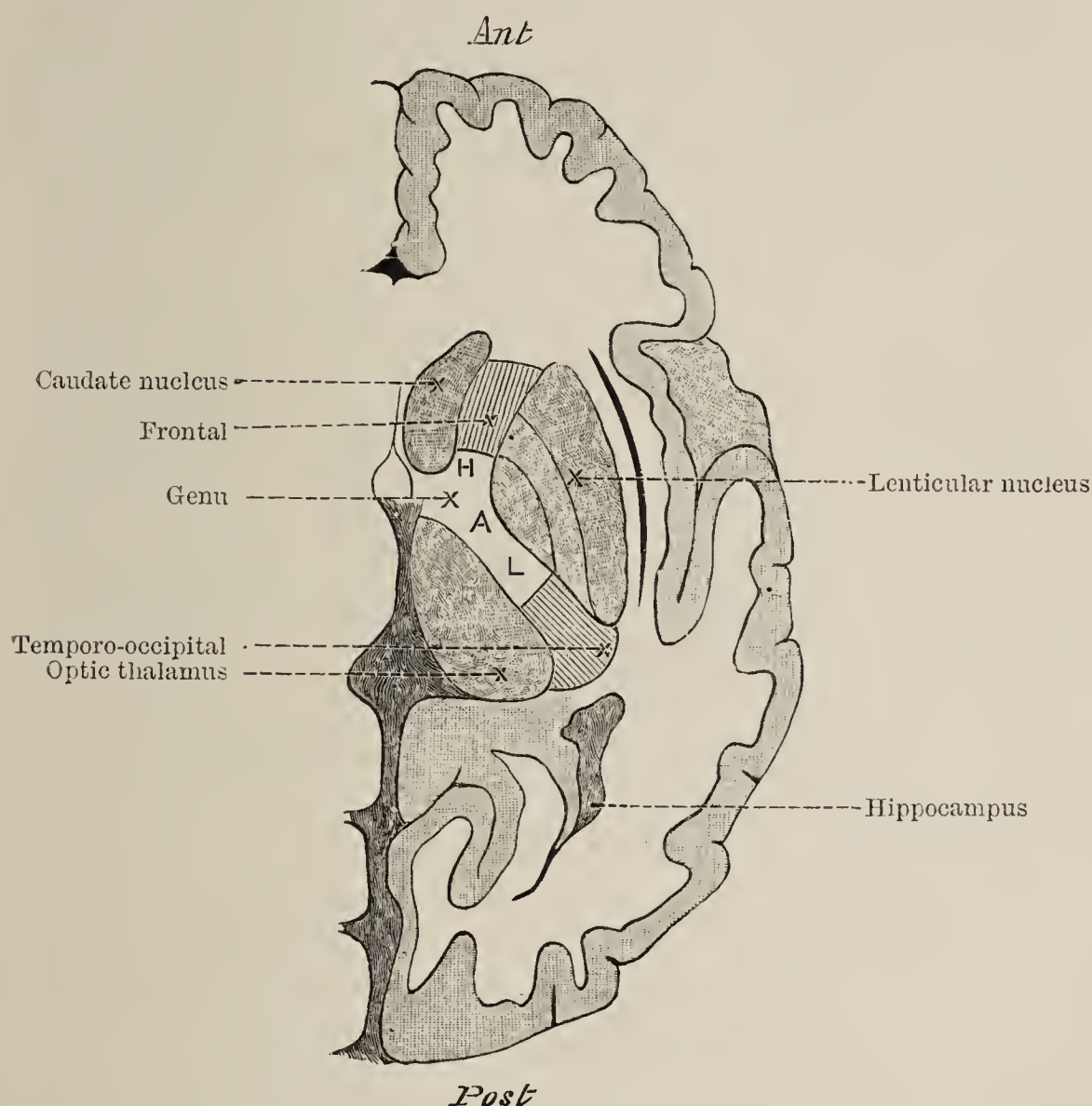


FIG. 282.—IDEAL HORIZONTAL SECTION THROUGH THE RIGHT HEMISPHERE AND BASAL GANGLIA.

To illustrate the position of the internal capsule taken in transverse section; H A L indicate the situations in the internal capsule of fibres governing the movements of the head, arm, and leg respectively. (After Charcot.)

the main channel of motion and of sensation. This main channel is continued as the two *crura cerebri*, which traverse the pons Varolii, and are continuous with the white columns of the bulb and cord. It is probable that this main stream gives and takes fibres to and from the ganglionic masses which it skirts or traverses at the base of the brain, but to what extent we cannot say, and it is important to realise that a large portion passes directly through, as a system of afferent and efferent channels, between



grey matter of cortex and grey matter of cord. The clearest testimony of this is afforded by the tract of descending degeneration that has been traced from cortex through corona radiata, internal capsule, crus cerebri, pons, anterior pyramids of bulb, to the lateral column of the opposite side (the crossed pyramidal tract), and to the anterior column of the same side (the direct pyramidal tract).

As regards localisation of motor and of sensory paths in the internal capsule and in the crus, pathological observations go to show that the posterior division of the internal capsule and the anterior or ventral portion of the crus (pes, crusta, or basis) are the main channels of motor and sensory impulses. It has further

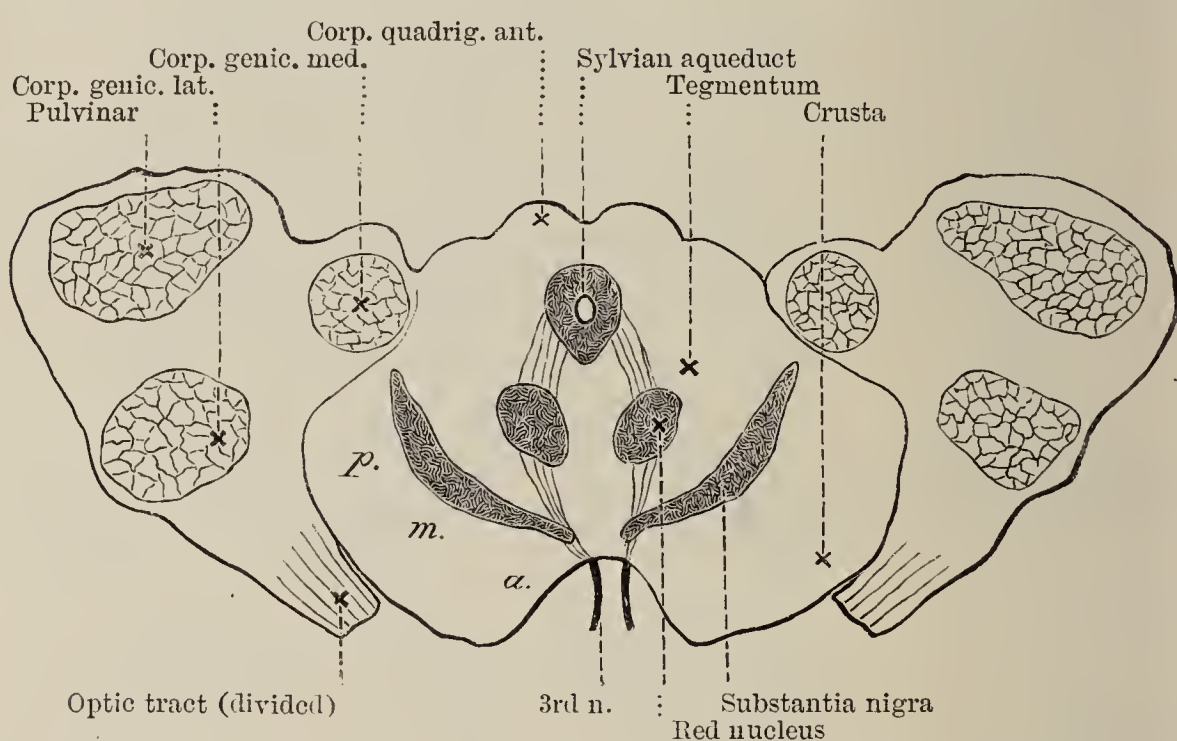


FIG. 283.—DIAGRAMMATIC TRANSVERSE SECTION THROUGH THE CRUS CEREBRI AND ANTERIOR CORPORA QUADRIGEMINA. (After Obersteiner.)

The letters *a. m. p.* on the pes or crusta signify portions occupied by fibres from the anterior, middle or Rolandic, and posterior regions of the cortex.

been observed that injury of the anterior two-thirds of the posterior division of the internal capsule is associated with hemiplegia; injury of the posterior third, with hemianæsthesia; injury of the whole, with hemiplegia *cum* hemianæsthesia. Descending degeneration from the frontal and occipital regions of the cortex are respectively anterior and posterior to the pyramidal area in the capsule and crus, but are not traceable beyond the pons; whereas the pyramidal degeneration from the Rolandic region extends down the spinal cord (p. 492).

In addition to (1) the vertical system of fibres above described, the corona radiata contains (2) a horizontal system of fibres between the two hemispheres, having their chief path through

the *corpus callosum*, and (3) numerous but scattered association fibres between the various convolutions. These last-named fibres are difficult to trace, least so, perhaps, in the gyrus fornicatus, in which there is a well-marked longitudinal system known as the cingulum. The vertical and horizontal fibres, derived respectively from the internal capsule and the corpus callosum, are also difficult to dissociate, being closely interwoven in their expansion through the corona radiata.

*Relation of cerebral development to intelligence.*—The brain, or, more precisely speaking, the cortex of the brain, is the organ of intelligent sensation and motion. Taken in the rough, the intellectual rank of animals bears some relation to the weight of the brain. Thus the average ratio between brain-weight and body-weight is in round numbers—

In fishes . . . . .	1 to 5,000
„ reptiles . . . . .	1 to 1,500
„ birds . . . . .	1 to 220
„ mammals . . . . .	1 to 180
„ ourang . . . . .	1 to 120
„ man . . . . .	1 to 50

But mere weight of brain is not a regular index of degree of intelligence in individual cases ; there are many exceptions to the general rule.

As regards man very similar considerations hold good, viz. taken in the rough, the brain-weight of distinguished men is above, while that of idiots is below, the average, and the brain-weight of civilised men is above that of savages. Yet there are many individual exceptions to the general rule. The average brain-weight is :

Of Europeans . . . . .	49 oz., or 1,390 grammes
„ negroes . . . . .	44 oz., or 1,250 grammes

As regards sex, the brain-weight cannot be appealed to in evidence of superiority on either side. The average male European brain weighs 49 oz.; the average female European brain weighs 44 oz.; but the average body-weights in the two sexes

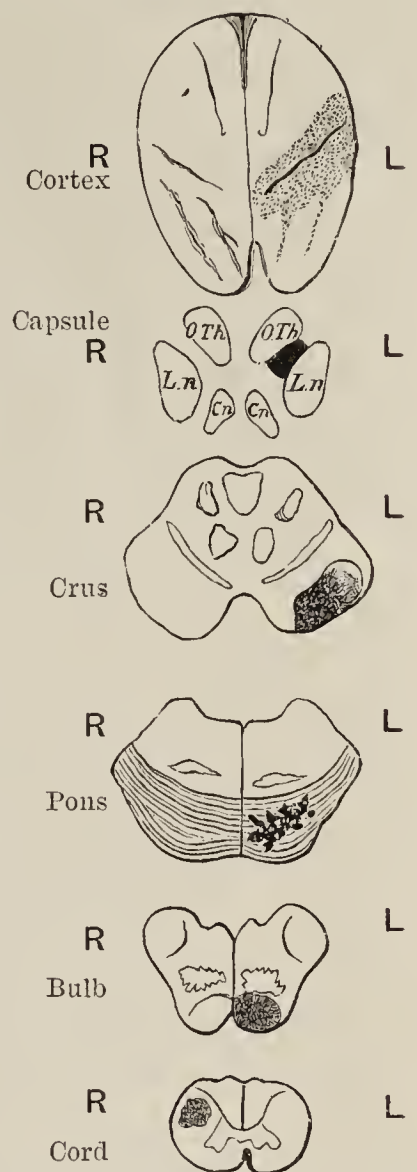


FIG. 284.—TO ILLUSTRATE THE PYRAMIDAL TRACT. (After Gowers.)



differ in about the same proportion. There is a more precise relation, though by no means an absolute one, between quality of brain and intelligence. The brain of an animal high in the scale is more complex than that of an animal low in the scale; higher in the scale, the hemispheres are relatively more developed, both as regards size and as regards the complication of their gyri. In man the hemispheres have their maximum development; they cover all the rest of the cranial contents; their surface-convolutions are of almost bewildering complexity. And now, taking a series of human brains, and comparing the convolutions of uncivilised and of civilised men, of men distinguished by abilities much above or much below the average, a general relation is traceable between complexity of surface and degree of intelligence. The basal ganglia are identical in the two cases, but the higher brain is more richly convoluted, its sulci are more numerous; the lower brain is simpler, its sulci are less numerous; the latter is practically a simplified diagram of the former; the effect of the difference is that the total area of grey matter is greater in the more highly than in the less highly organised brain. This is all we know with regard to the relation between quality of organ and quality of function; finer and more impalpable relations doubtless exist, but have not been demonstrated; the very absence of anatomical difference in the brains of average and of exceptionally able men leaves us, however, free to think, and justified in believing, that beyond quantity of grey matter there are differences in its quality. Perhaps its cells are more numerous, perhaps these more numerous cells are of better quality in the brains of the exceptionally able; but no microscopical or chemical proof has been given for or against such suppositions, still less is there any physical evidence available to distinguish the brain matter of 'good' men from that of 'bad' men.

Before entering upon the detailed consideration of the cortex cerebri, we may pass in review such knowledge as we possess regarding the basal ganglia and the cerebellum. The term basal ganglia includes the corpora striata and the optic thalami, and may be taken to embrace the less prominent ganglionic eminences known as the corpora quadrigemina (= the optic lobes of the lower vertebrates) and the corpora geniculata.

**The basal ganglia. The cerebellum.**—Our definite physiological knowledge concerning the part played by the chief basal ganglia (*optic thalamus, corpus striatum*) is extremely scanty. They form

prominent and imposing masses of grey matter at the isthmus of the brain, but what part they may play in the elaboration of nerve-impulses we do not know. From their anatomical situation they were formerly characterised as 'ganglia of interruption,' on the motor and on the sensory tracts respectively, but this conjecture—at least as regards the corpus striatum—has been more and more weakened by additional pathological information; and as to their embryonic origin, the chief fact, whatever its significance may be, is that they are derived from different cerebral vesicles of the fore-brain—the corpus striatum from the first vesicle, *i.e.* that of the hemispheres (prosencephalon), and the thalamus from the second vesicle (thalamencephalon). The chief motor tract merely skirts the ganglionic matter of the corpus striatum, without forming connections with its cells.

Sensory tracts undergoing 'ascending' degeneration have been traced to the thalamus; and after extirpation of the cortex the thalamus suffers degeneration—or rather atrophy.<sup>1</sup> The balance of evidence points to the thalamus as a link in the sensory chain, to the corpus striatum as a terminal organ.

Anatomical connections and lines of degeneration are indeed the chief source of such knowledge as we possess concerning the possible functional relations of these and other basal ganglia—corpora geniculata and corpora quadrigemina. The optic nerve is anatomically connected with the external geniculate and anterior quadrigeminal bodies and with the pulvinaria of the optic thalamus; through these three stations, more especially the first—

<sup>1</sup> The atrophy or deficient generation of nerve-tracts and ganglia, often spoken of as 'degeneration,' is not to be confounded with Wallerian degeneration (p. 356); the latter is the rapid and gross effect of active malnutrition, the former (to which reference is made in the text) is a gradual effect of functional disuse. In the case of an enucleated eyeball, there is Wallerian degeneration of the optic nerve and tracts (direct and crossed portions) as far as the external geniculates, 'degeneration by disuse' of and beyond these ganglia (chiefly on the opposite side). An early stage of degeneration by disuse in fibres and cells *above* the point at which a nerve has been cut has recently been described by Durante and by Marinesco; clumps of stainable material, recognisable in the normal cell and fibre under a high power of the microscope, breaking up and disappearing within a day or two after the operation. Degeneration by disuse is also observed in the nerves and nerve-centres belonging to an amputated limb, and is sometimes referred to as retrograde degeneration.

Regarding the nerve-cell with its processes (inclusive of the axis cylinder of a fibre) as one individual (or 'neuron') we are to recognise that *both* portions into which it is divided when a nerve is cut across, undergo degeneration—rapid Wallerian degeneration of the distal part, and gradual 'retrograde' atrophy of the proximal part and of the cell body.



named, it is connected with the occipital cortex by the optic radiation of Gratiolet. All these parts, especially in young animals, have been found atrophied as a result of destruction of the opposite eyeball; on the other hand, destruction of the occipital cortex has been found to produce atrophy of the optic radiation, external geniculate body, optic tract and chiasma down to the retina itself (Monakow). From which we may at least conclude that the external (= lateral) geniculate body is a link in the chain of visual sensation.

On a similar footing, in view of their anatomical connections with the cochlear nerve, we may admit that the posterior quadrigeminal and internal geniculate are in the path of auditory sensation.

As to smell, the fact that the olfactory bulb is anatomically connected with the hippocampal gyrus and the amygdaloid nucleus needs no further comment.

With regard to the part played by the cerebellum, our position from a physiological point of view is little better. A bulky organ cannot but be of importance; anatomically, the cerebellum presents features that seem to promise great functional significance; it is connected by definite tracts with the cerebral hemispheres and with the spinal cord, it exhibits on section a more definitely regular structure than any other part of the nervous system, and histologically the disposition and regularity of the large cells of Purkinje seem to say that they play a definite and easily-discoverable part in the control of the body. Yet we have not during the last fifty years added anything to, but have made

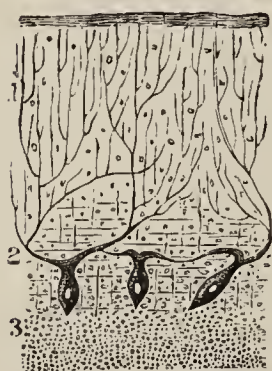


FIG. 285.—TRANSVERSE SECTION THROUGH THE CORTEX CEREBELLI.  $\times 50$  diam. (Obersteiner.)

reservations to and actual subtractions from, the conclusion of Flourens in 1831, that the property of the cerebellum consists in co-ordinating the movements incited from other nervous centres. Now, as then, the one definite function that we assign to the cerebellum is muscular co-ordination, the one prominent symptom of cerebellar disease is a staggering gait, and the only experimental effects of destruction or of excitation of different parts of the organ are forced movements; but these,

as we shall see, are producible by lesions of many other parts of the brain. Clinically, the cerebellum is supposed to play an important part in the government of the muscles and the maintenance of their 'tone'; but to the experimental physiologist

it is an organ of disappointment. There is no paralysis after destruction of the whole cerebellum. There is no loss of intelligence. There is no evidence in favour of the conjecture that the cerebellum has anything to do with a genetic sense. There is nothing to prove that the vertigo produced by electrical currents applied to the back of the head is cerebellar; it may equally well be due to excitation of the semicircular canals.

Of the three principal divisions forming the cerebellum of man, the median lobe, or vernis—which in birds forms the entire organ—is, according to clinical observations, the most important, and, when damaged, most usually productive of characteristic signs. Lesions or excitation of this lobe, to be of any effect, must be unilateral, and the chief facts that have been observed are as follows:—Unilateral injury causes movements of rotation of the body *towards* the side of injury (*i.e.* clock-wise if the injury is on the right), and inclination to fall to the opposite side. Unilateral excitation causes muscles to contract on the same side, and produces conjugate movements of the eyes to that side. After unilateral ablation, the excitability of the cerebral cortex is greater on the opposite than on the same side (Russell). Dissections and degenerations indicate that the cerebro-cerebellar connection is a crossed one, chiefly *via* the internal capsule and red nucleus of one side, and superior cerebellar peduncle of the opposite side, while the spino-cerebellar connection in the main is uncrossed *via* the restiform and cerebellar tracts. In this scarcity of significant experimental results we are thrown back upon purely hypothetical inferences from anatomical connections; the cerebellum is a nerve-centre, therefore an organ of return of action; it has wide-spreading anatomical connections, probably afferent as well as efferent, with the cortex, with the basal ganglia, with the spinal bulb and cord, and through this first line of connections, with the whole receptive and motor periphery; when it is destroyed, muscular action is disorderly. Therefore cerebellar action is an essential ingredient in the reflex guidance of muscular action.

*Forced movements.*—We may take this opportunity of giving a short account of forced movements. Their characteristic is an overmastering action of the muscles of one side, or of large groups of ‘synergic’<sup>1</sup> muscles acting in locomotion, in consequence of which the body twists round its long axis, or progresses

<sup>1</sup> ‘Synergic’ muscles are such as act together in habitual movements effected by the associated action of several muscles.



in a circle of small radius (circus movements) ; sometimes the excessive action is such as to cause an animal to precipitate itself straight forwards, and to turn somersaults in its progress, sometimes the tendency is to recede. It is probable that the central condition is one of excitation rather than paralysis, for hemiplegia is not attended by forced movements. With regard to the possible seat of excitation, we may recognise (1) that it is not in the cortex, the excitation of which, as we shall see, gives rise to epileptic movements ; (2) that it may be due to lesions of the cerebellum, of the pons, of the crus cerebri, or of the basal ganglia ; (3) that it may originate from the semicircular canals.

We may in some degree realise the nature of a forced movement by applying a constant current through two electrodes fixed to the back of the head. The head and body feel as if bent towards the anode, and in spite of our voluntary resistance the inclination becomes irresistible ; the body bends and falls to that side, and the accompanying sensation is that of being dragged down ; the eyes undergo rhythmic conjugate jerks (nystagmus) towards the side of the kathode. We have experienced a forced movement, and although under these conditions we may be in doubt whether the effect is by an uncontrollable excitation of the cerebellum, or of the semicircular canals, we may, from observations on birds, see reason to prefer the view that the semicircular canals rather than the cerebellum are the effective agents. Ewald has shown on pigeons that such forced movements produced by galvanism on the intact bird, are no longer produced after destruction of the semicircular canals, and are but little interfered with by destruction of the cerebellum. Pollak, on applying this galvanic test to deaf-mutes, found that no less than 30 per cent. gave no responsive movements of the head and eyes ; presumably in such cases the semicircular canals were destroyed, and the test has an obvious possible application in precise diagnosis.

Vertigo, or giddiness, furnishes a good illustration of the character of forced movements ; subjectively, *i.e.* as felt by the subject, the prominent symptoms are distorted and exaggerated sensations of external objects, and uncontrollable muscular inclinations ; objectively, *i.e.* as seen by an observer, the prominent sign is staggering, *i.e.* uncoördinate forced movements.

## THE CORTEX OF THE BRAIN

The cortex, or grey mantle of the brain, is divisible into the following regions: frontal, parietal, occipital, temporal, mesial, and insular. The frontal region comprises the superior, middle and inferior frontal convolutions, and the ascending frontal or præcentral convolution, which is situated in front of the fissure of Rolando, or central sulcus. The parietal region comprises the superior and inferior parietal lobules, separated by the intra-parietal fissure, and the ascending parietal, or post-central convolution, situated behind the fissure of Rolando; in the inferior parietal lobule are to be recognised the supra-marginal gyrus at the posterior extremity of the Sylvian fissure, and the angular gyrus at the posterior extremity of the first temporal fissure. Three occipital convolutions are distinguished in the occipital region, and three temporal convolutions in the temporal region. The mesial aspect of the cortex, exposed by separation of the two hemispheres, includes the marginal convolution and gyrus fornicatus, separated by the calloso-marginal fissure, and the hippocampal and uncinate gyri; the gyrus fornicatus is continuous with the gyrus hippocampi, forming with it the 'grand lobe limbique' of Broca, which should not be confused with 'Broca's convolution,' *i.e.* the third left frontal. The insular region, or island of Reil, is exposed by separating the lips of the Sylvian fissure; it is formed by a group of short convolutions occupying the lower surface of the frontal lobe (see figs. 289, 290).

*Microscopically*, the grey matter forming the cortex of the brain consists of nerve-cells, nerve-fibres and neuroglia disposed in layers that are more or less distinguishable from each other. The stratification is best marked in the hippocampal region, which is characterised by a very regular layer of large nerve-cells; in other parts it is far less definite, so that it is almost impossible to identify each of the five layers of classical description (Meynert). A section of grey matter from a 'motor' convolution is, however, to some extent distinguishable by the prominence and number of large arrow-head cells (30 to 40 to 100 ) in the deeper part of the so-called third layer; these are sometimes characterised as 'motor,' and are considered to be the analogues of motor cells in the anterior cornua of the cord and of Purkinje's cells in the cerebellar cortex. It nowise follows, however, because a nerve-cell is large, that therefore it is motor, and as regards the various cells above alluded to, those of the



anterior cornua are the only ones whose claim to the term motor may be regarded as established upon sufficient data. The

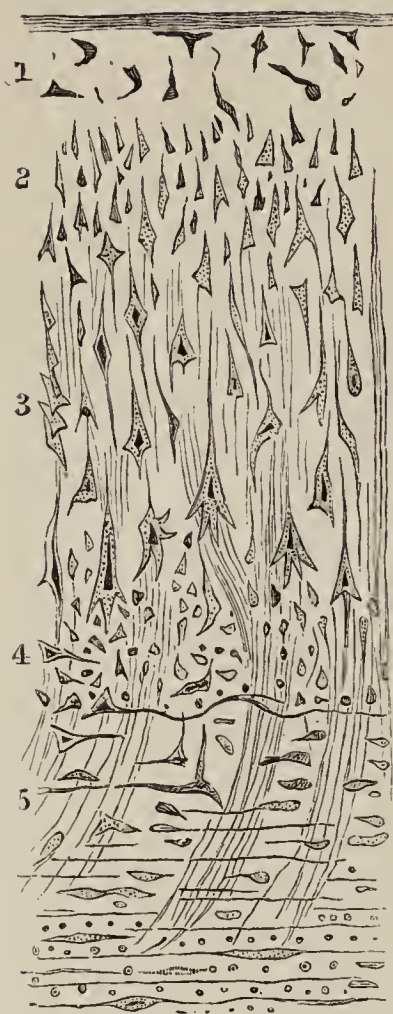


FIG. 286.—TRANSVERSE SECTION THROUGH THE CORTIX CEREBRI. FIVE-LAMINATED OR MOTOR TYPE  $\times 50$  diam. (Meynert.)

'motor' cells of the cortex have been so characterised because they are a prominent feature in the experimentally excitable area of the brain; other large cells, whether in the hippocampus or in the cerebellum, are not qualified by any such designation. And, as a matter of fact, it happens that the large-cell layers are most definite in the very regions (hippocampus, cerebellum) the functions of which are most indefinite, and in all probability not motor. The supposition that the 'motor' cells of large muscles, or of 'large movements,' or connected with long nerve-fibres, are 'large' as compared with 'motor' cells of small muscles, or of 'small movements,' or of short nerve-fibres, is a speculation as yet unsupported by any actual proof. The total number of nerve-cells in the human brain has been estimated at 2,000 millions.

*The vascular supply of the brain.*—The arterial system of the brain is peculiar in two main particulars: (1) the large arteries by which it is supplied—two internal carotids and two vertebrals—form a free anastomosis at the base of the brain, known as the circle of Willis; (2) the branches that are given off from the circle of Willis, supplying the basal ganglia and the cortex, run their course to their several areas of distribution with but little anastomosis. It follows as a consequence of this disposition of the vessels, that the circulation in the brain suffers little or no change by obstruction on the cardiac side of the circle of Willis, and that, on the other hand, obstruction of a vessel beyond the circle of Willis causes damage which is confined to the area served by the obstructed vessel. The first statement is borne out by the experimental fact that, *e.g.* on dogs, it is not until all the four vessels leading to the circle of Willis are tied that symptoms of arrested cerebral circulation appear. The second statement is most clearly exemplified by the well-characterised group of clinical symptoms that are caused when the left middle cerebral artery is blocked by an embolus. Considered from the point of

view of its vascular supply, the cortex of the brain is divisible into three regions, supplied by the anterior, middle, and posterior cerebral arteries respectively. The cortical territory fed by the *middle cerebral*, or *Sylvian artery*, which is the direct continuation of the internal carotid, is physiologically and pathologically the most important; it includes the convolutions in the vicinity of the fissure of Rolando, and those of the island of Reil, *i.e.* the so-called motor area and speech-centre. If the Sylvian artery is blocked (and in the great majority of cases this happens on the left side), the consequences are motor paralysis of the right side and loss of speech, *i.e.* in clinical language, right hemiplegia and motor aphasia. The posterior cerebral artery, more particularly its *occipital* or *calcarine* branch, supplies the area concerned in visual perception.

In addition to the cortical system formed by branches of the three cerebral arteries, other branches from them are supplied to the basal ganglia, and form the ganglionic system of arteries. Those given off by the middle cerebral are of greatest pathological importance, as being the vessels most apt to give rise to hæmorrhage; one branch in particular, the *lenticulo-striate*, has been called the '*artery of cerebral hæmorrhage*' (Charcot), because it is most liable to give way; the hæmorrhage derived from it may compress or rupture the internal capsule, and so cause hemiplegia. As regards the ultimate distribution of the cerebral capillaries, the most noteworthy point is, that the network is much closer and richer in grey matter than in white matter; the former is physiologically the more active, and requires to be more abundantly nourished with blood; a similar difference holds good in the case of the spinal cord, in well-injected sections of which the closer vascular network of the grey matter defines the pattern of the cornua in the midst of the scantily supplied white matter.

The return of blood from the cortex is chiefly effected by a superficial system of veins opening into the superior longitudinal sinus, and the fact that, whereas the current in the sinus is directed from before backwards, the veins opening into it are directed from below upwards, has been remarked upon as promoting venous stagnation, giving rise, therefore, to a peculiar liability to thrombosis.

It was formerly a debated question whether or no the amount of blood can vary within the cranium; taking into account that the cerebral ventricles are in communication with the sub-arach-



noid space and filled with a serous fluid, there can be no doubt that more or less blood may enter the cranium, expressing more or less serous fluid into the spinal canal. The sub-arachnoid and intra-ventricular fluid is, indeed, of great mechanical importance, forming as it does a bath within which the cerebro-spinal mass is protected from accidental shocks and sudden pressure.

**The functions of the brain.**—That the brain is the organ of intelligent sensation and motion is proved by the facts of comparative anatomy already alluded to, and by common experience. The same proposition is established by clinico-pathological facts, and by the study of animals after removal of a hemisphere or of the cortex. Experimentally, we learn that after removal of the cortex an intelligent animal is reduced to the state of a non-intelligent automaton, responding indeed to stimuli, internal as well as external, but failing to interpret the significance of present events in accordance with bygone experience. A brainless dog is stupid; he may see a bone in front of his eyes without showing sign that he knows the meaning of a bone, or the use to which it may be put; he may hear the crack of a whip, but he no longer shows sign of fear, for he does not remember its sting; his former

purposeful behaviour has entirely disappeared: in short, he has lost memory and judgment.

From clinico-pathological observations, as well as from experiment, we learn, moreover, that each hemisphere is the controlling organ of the opposite side of the body. Clinical cases of ordinary hemiplegia are every day found to be due to unilateral disease in the opposite hemisphere. Experimental extirpation or stimulation on one side of the brain is found to produce paralysis or muscular contractions of the opposite side of the body. These facts are, of course, referable to the crossing of motor and of sensory paths in the bulb and

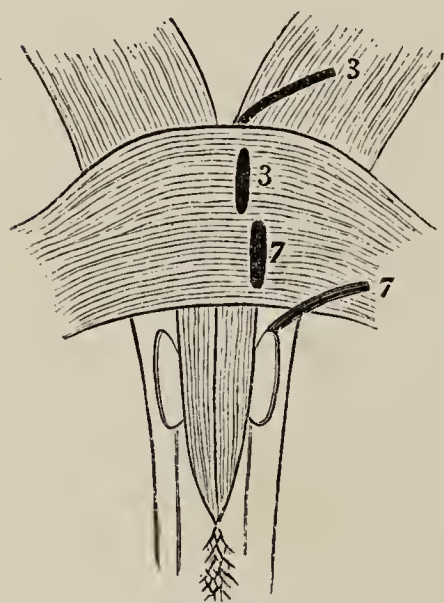


FIG. 287.—DIAGRAM OF THE ORURA, PONS, AND BULB.

To illustrate the mechanism of crossed paralysis; the figures 3, 7, point to the nerves and nuclei commonly involved.

cord. Typical common hemiplegia is limited to one side, but not infrequently cases of *crossed paralysis* occur, in which the arm and leg of one side and the face on the other side are paralysed, and it is found post-mortem that the combination is due to

lesions on one side of the brain involving nerves or nerve-nuclei at the base of the brain, which are controlled from the cortex of the opposite side. A tumour in the vicinity of the pons, for instance, might interrupt the left pyramidal tract, and at the same time the left facial nerve or its bulbar nucleus; under these circumstances, right hemiplegia of the limbs in combination with paralysis of the left side of the face would be produced, and the face would be drawn towards the paralysed side of the body, instead of from that side, as in a case of common hemiplegia.

The fundamental notion that the muscles of one side of the body are connected with the opposite side of the brain requires to be supplemented by certain other considerations; it cannot be doubted that each side of the brain mainly governs muscles—or, more properly speaking—movements of the opposite side of the body, especially in their unilateral and most highly specialised modes of action; but it is no less certain that from one side of the brain muscles of both sides of the body may be governed, this being especially the case in their bilateral and more automatic action. This principle is exemplified in the everyday clinical phenomena of hemiplegia; it is particularly striking in conjugate movements, as in the case of the head and eyes, and in bilateral movements, as those of the respiratory and laryngeal muscles; it is also seen in the effect of experimental stimulation of one side of the cortex, a weak stimulus giving movement only on the opposite side, but a strong stimulus giving movement also on the same side of the body, though with less energy and with a greater lost time than on the opposite side.

In a typical case of hemiplegia all the muscles of one side of the body are not equally affected; the muscles of the extremities are most completely paralysed, those of the trunk not at all, or only partially on both sides. More generally expressed, the muscles that suffer most are muscles with unilateral action, engaged upon specialised movements; those which suffer least are muscles that are bilaterally associated in their actions. This bilateral association postulates communication across the middle line between symmetrical centres of the two sides of the body. The communication may be effected by commissural fibres between the cortical centres, or more probably by commissural fibres between bulbo-spinal nuclei, for it has been shown experimentally that excitation of one side of the cortex can produce movements on both sides of the body after the cerebral commissures have



been divided, or after the opposite hemisphere has been destroyed. In unilateral convulsion (*e.g.* hemichorea) there is a similar but reversed inequality in the distribution of spasm on the two sides of the body; in the case of muscles of unilateral action the convulsions may be limited to one side, but in that of muscles habitually associated in bilateral action the convulsions always affect both sides. Moreover, it is possible that a muscle may be paralysed as regards its participation in a voluntary highly specialised manœuvre, while remaining efficient as regards its participation in reflex or in automatic action.

From all these facts two important consequences are to be drawn: (1) that each hemisphere, while mainly governing the opposite side, also takes part in the control of the same side; *i.e.* from any given cortical focus not only crossed fibres must exist, but also direct (or more probably twice-crossed) fibres; (2) that particular movements rather than particular muscles are associated with cortical motor foci. The significance of this second statement will be better appreciated after the doctrine of localisation has been considered.

A remarkable feature in the history of a paralysis, especially one of cortical origin, is that the loss of motility gradually diminishes, as if the cortical blot were encroached upon from its periphery. A paralysis or a paresis in consequence of a central lesion, may theoretically be due to a cessation, and therefore absence of action, or to an active arrest of action. The ordinary and popular view is, that it is always due to an absence of action, and no doubt absence of action does occur in connection with destruction of nerve tissue. But we must also recognise that destroyed tissue is surrounded by excited tissue, and admit, with Brown-Séquard and Goltz, that a paralysis or paresis may in many instances be due to an active arrest of action—*i.e.* an *inhibition* originating in such exciting tissue. The variability and the curability of many paralytic phenomena are intelligible to us on this basis, *viz.* as due to the functional fluctuations of disturbed tissue; and it is important to recognise that such disturbed tissue may give rise to paralysis by 'inhibitory action,' as well as to convulsion by 'dynamogenic action' (Brown-Séquard).

Finally, it should be mentioned as a possibility—or rather as a probability—that over and above the gradual effacement of acute functional effects, there is a gradual assumption of control over paralysed parts by associated uninjured cortex. This is particularly probable in the case of animals, and in that of man

when lesions are of limited extent (p. 543); it is also believed to have occurred in cases of aphasia (p. 549).

*Conjugate movements of the eyes, head, and neck.*—We have learned in the section on Vision that the movements of the two eyes are exactly equal and parallel for different directions of distant vision; both eyes are turned to the right or to the left, up or down, so that the object fixed gives images on corresponding parts of both retinæ. In movements directly upwards or downwards muscles of the same name in each eye are associated in action; but in lateral movements the association is asymmetrical, *e.g.* the external rectus of one eye acts with the internal rectus of the other; and the peculiarity of this associated action seems still more striking

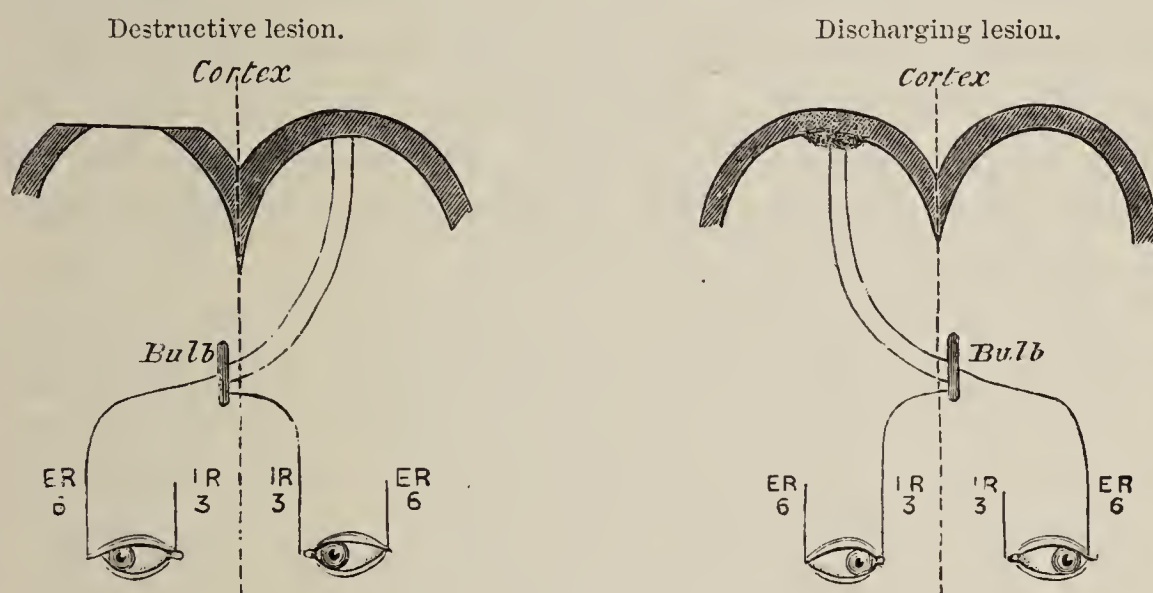


FIG. 288.

Diagram to illustrate conjugate deviation of the eyes—that is, *away from* a discharging lesion, *towards* a destructive lesion. In the diagram the eyes are supposed to be turned towards the left by a discharge from the right side of the cortex; to the right, by unbalanced action of the left side of the cortex if the right side is destroyed. In both cases the head rotates to the same side as the eyes.

when it is remembered that the external rectus is supplied by the sixth nerve, while the internal rectus is supplied by the third. A similar, if less striking, association of asymmetrical muscles on the two sides occurs in the rotation of the head and neck, which are turned to the right by the right inferior oblique and left sterno-mastoid muscles, to the left by the left inferior oblique and right sterno-mastoid. In looking to the right we contract the right external and left internal rectus, *i.e.* impulses pass through the right sixth nerve and the left third nerve, possibly from the left and from the right side respectively of the motor cortex, but more probably from only the left motor cortex, in which case we must suppose that certain nerve-fibres cross twice, once between cortex and bulbar nucleus, and a second time between nucleus and nerve-



termination. That the mechanism is actually of this nature is indicated by the effects of cortical convulsions and of cortical hemiplegia upon the position of the eyes. Unilateral convulsions of cortical origin are accompanied by rotation of the head and eyes towards the convulsed side, *i.e.* away from the cerebral lesion. Thus a discharging lesion of the right motor cortex causes convulsions of the left side of the body, with rotation of the eyes to the left. This is a '*conjugate deviation*,' and the fact is peculiar, inasmuch as we have discharge from the right side of the brain, along the left sixth nerve (external rectus) and the right third nerve (internal rectus).

A destructive lesion of the right motor cortex causes paralysis of the left side of the body, with rotation of the eyes to the right. The peculiarity in this case is, that there is cessation of action along the left sixth nerve (external rectus) and the right third nerve (internal rectus), the deviation of the eyes to the right being caused by the unbalanced action of the muscles which rotate the eyes to the right.

Movements of the eyes to the right or left are habitually associated with rotation of the head and neck in the same direction. The head is turned to the right by the left sterno-mastoid and right inferior obliquus capitis, and *vice versa*; these, then, are conjugate muscles, acting like the eye muscles above considered, and in conjugate action with them. This conjugation is preserved in the spasmodic or paralytic deviations caused by cerebral lesions, the head and neck being always turned in the same direction as the eyes.

#### CORTICAL LOCALISATION

The physiology of the brain has, since the year 1870, been engrossed by the one question of localisation. Is there a localisation of different functions in different parts of the brain, are its different parts concerned with different offices, is there a division of cerebral labour? or does every part subserve every office, can any office be administered by any part? To these questions it may be answered at once that the balance of evidence is in favour of localisation, but that the counter-evidence has shown that localisation is not sharply defined, but blurred. The state of doctrine may be made plain by an analogy. The various activities making up the business of the brain do not take place all over its surface, as in a country without towns or

villages, where all kinds of industry go on in every hut or tent ; nor are the different activities absolutely restricted to certain spots, as if in walled towns. The brain cortex is not comparable with either of these extreme cases ; its territory must be recognised as possessing towns with special industries, but towns with straggling and overlapping suburbs, and industries that are, indeed, predominant each in a given centre, but not exclusive of all other industries in that centre, nor excluded from other centres in which other industries predominate. In this qualified sense localisation of function in different parts of the brain must be considered to be established.

The chief facts by which the doctrine of localisation has been formed have been :—

(1) The proof given by Fritsch and Hitzig in 1870 that the cortex of the brain responds to experimental stimulation, and that the excitation of certain different spots provokes certain different muscular movements ; (2) the consequent detailed study—by Ferrier, Munk, Schäfer, Horsley, Exner, Nothnagel, Beevor, Goltz, Franc, Heidenhain, Luciani, Tamburini, Seppilli, Löb, and many others—of the relation between cortical areae, destroyed or stimulated, and resulting muscular paralysis or spasm ; (3) the scientific clinical study of epilepsy by Hughlings Jackson, completed by post-mortem recognition of the seats of lesion ; (4) the association found to exist between aphasia and lesion of the left third frontal convolution.

*The cortex of the brain is experimentally excitable.* This simple proposition was established by Fritsch and Hitzig in contradiction of the heretofore universally accepted statement by Flourens, Magendie, Longet, and the older physiologists, to the effect that the surface of the brain may be lacerated, pricked, burnt, mechanically, chemically or electrically excited, without provoking muscular contraction in any part of the body. Fritsch and Hitzig used dogs, exposed the surface of an entire hemisphere, and tested it by interruptions of the constant current and by induced currents. They found the lateral and posterior parts of the cortex to be non-excitabile, but in the frontal region they found certain spots, stimulation of which produced combined movements of the anterior or of the posterior extremities, of the neck, and of the face on the opposite side of the body ; they found that on closing or reversing the constant current, the anode was more effectual than the kathode, and they observed epileptic attacks following repeated



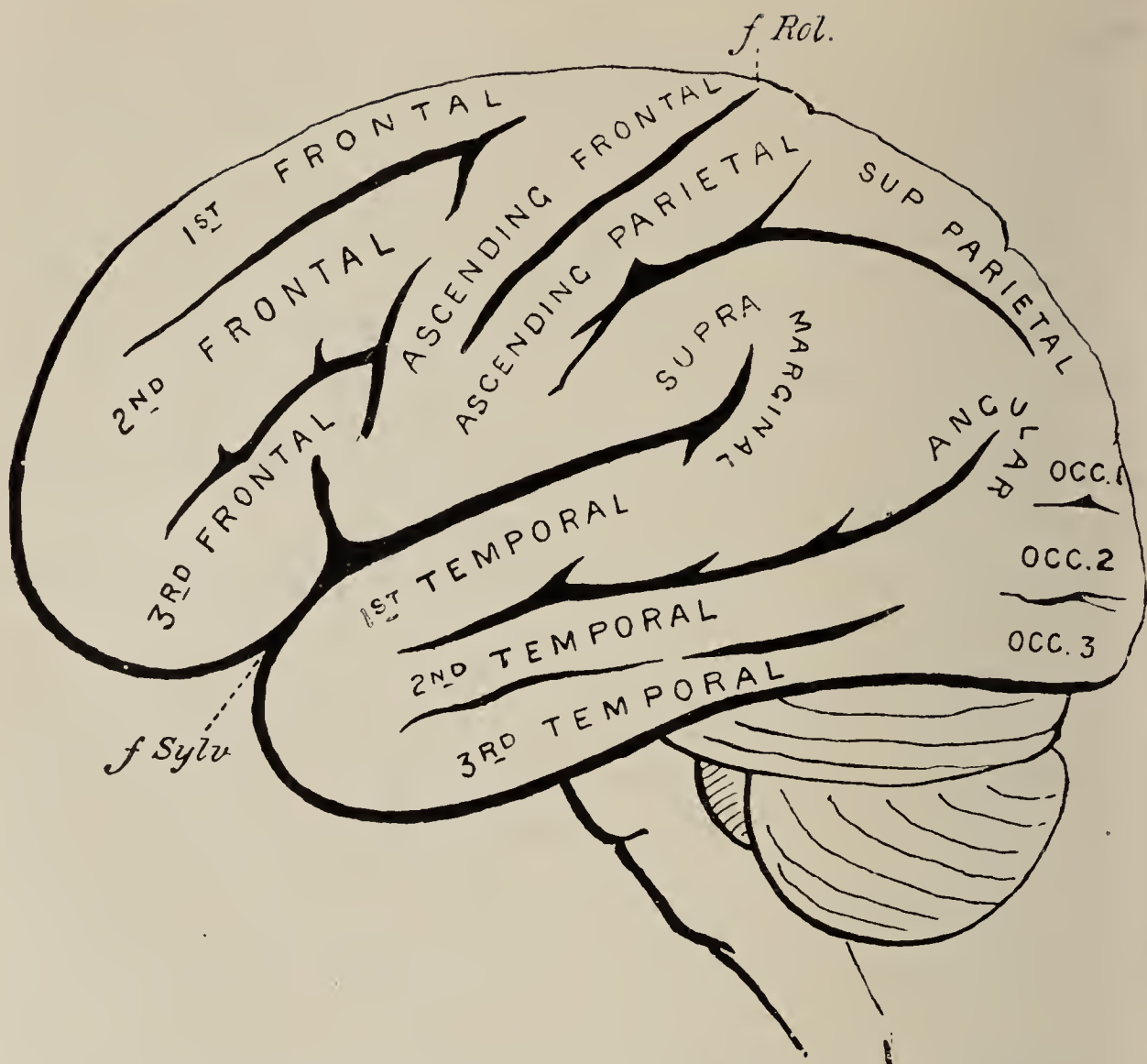


FIG. 289.—HUMAN BRAIN ; LATERAL ASPECT OF LEFT HEMISPHERE. (After Ecker.)

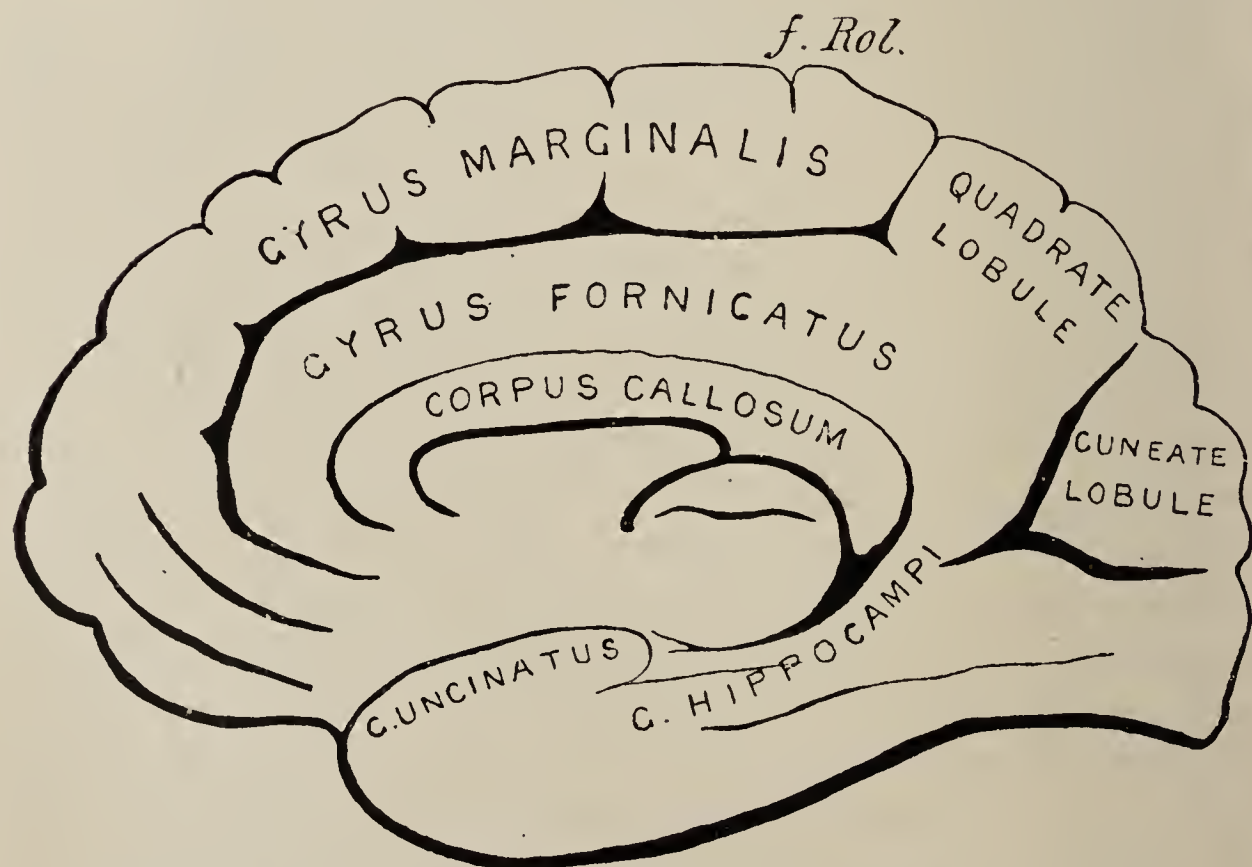


FIG. 290.—HUMAN BRAIN ; MESIAL ASPECT OF RIGHT HEMISPHERE. (After Ecker.)

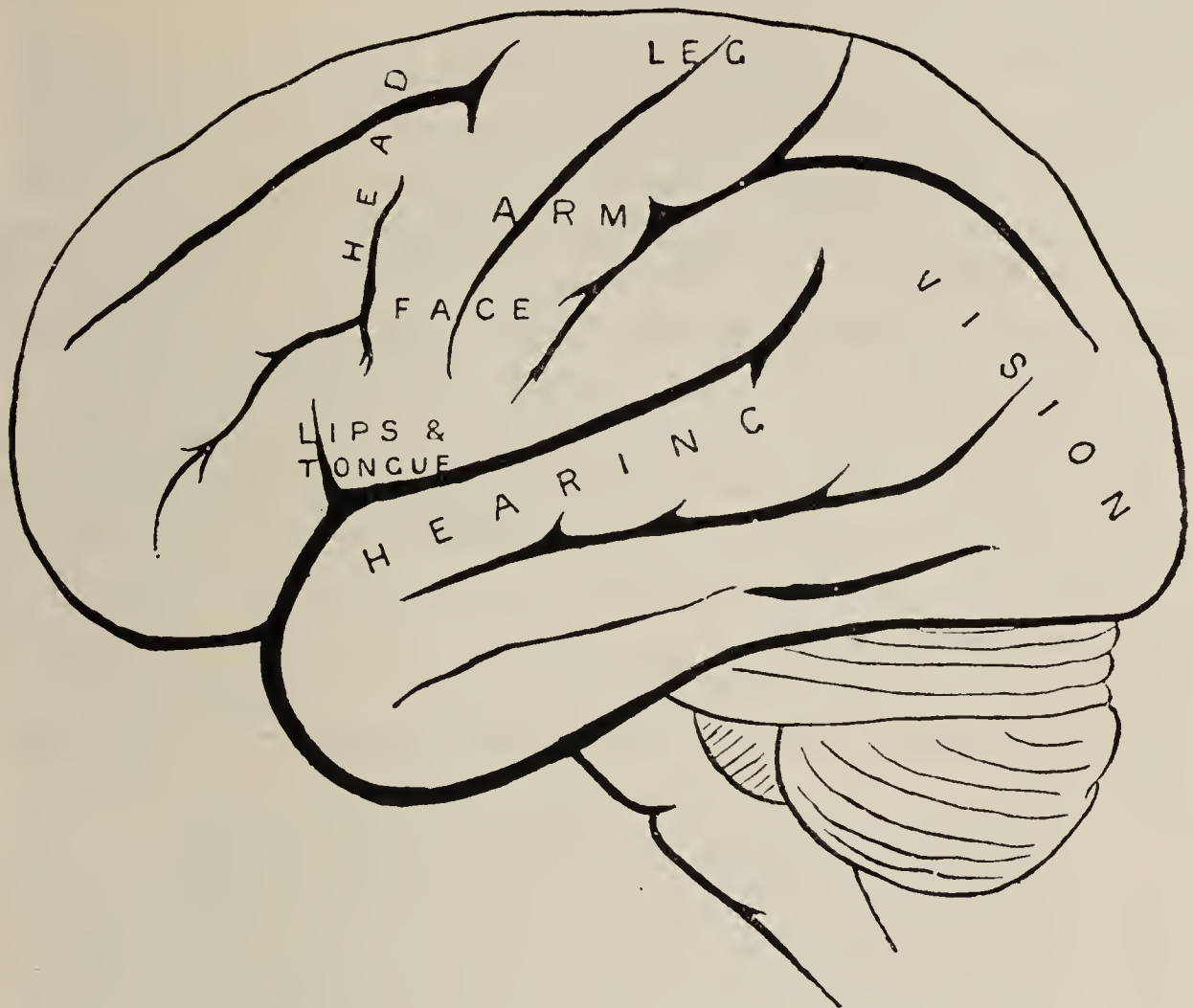


FIG. 291.—HUMAN BRAIN ; LATERAL ASPECT OF LEFT HEMISPHERE. TO ILLUSTRATE LOCALISATION OF FUNCTION.

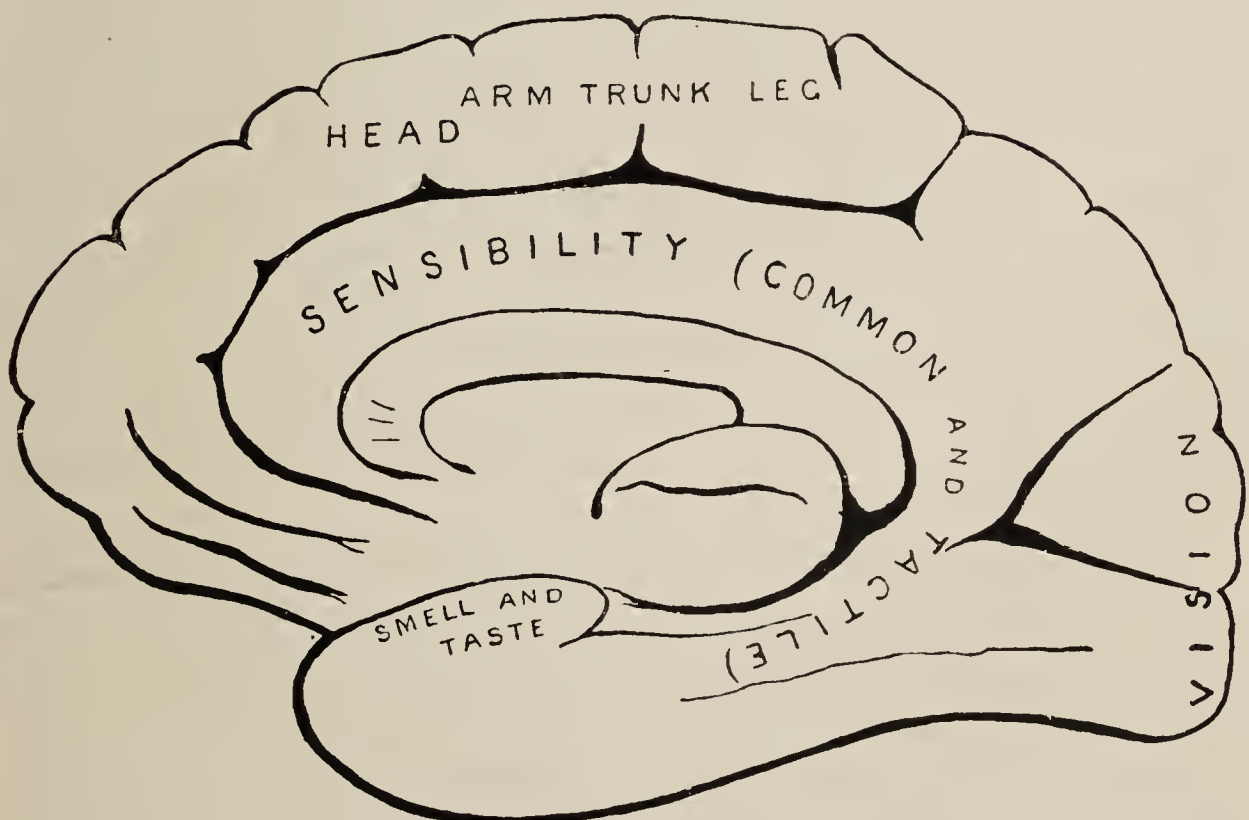


FIG. 292.—HUMAN BRAIN ; MESIAL ASPECT OF RIGHT HEMISPHERE. TO ILLUSTRATE CORTICAL LOCALISATION OF FUNCTION.



stimulation, and originating in spasms of the previously excited muscles.

*Objections.*—The electrical excitability of the cortex, the key-stone of the doctrine of localisation, was not at once admitted as proved. It was objected that the diffusion of currents in the brain-substance is such that stimuli cannot be limited to given areas, and that the effects are due to the excitation of the subjacent white matter or of the basal ganglia. To this objection the answers are: (1) No doubt the corona radiata is excitable; a localisation of effects is in fact demonstrable by excitation of different bundles from the cortex, as well as of different parts of the cortex itself; cortical stimulation remains without effect if the cortex is separated by an incision parallel to the surface, and replaced *in situ*; electrical diffusion to deeper parts is unaltered, but the physiological connection between cortex and base is interrupted. (2) Localised reactions are obtainable by mechanical stimulation of the Rolandic area. (3) Comparison of the 'lost times' in the two cases, viz. excitation of cortex and of subjacent white matter, reveals the fact that time is occupied in the physiological transmission of stimuli through the cortex; Franc and Pitres found, for example, on the dog, that the contraction of the front paw commenced .065 sec. after a cortical stimulus, .045 sec. after a stimulus to the subjacent white matter; *i.e.* they obtained .02 sec. as the time-value of cortical delay. These results were confirmed by Bubnoff and Heidenhain, and supplemented by a still more convincing proof of cortical excitability. They showed that the muscular contraction obtained by cortical excitation of a morphinised dog follows after a much longer interval, and is much more prolonged than that obtained immediately afterwards by sub-cortical excitation. (4) Confirmatory proof of the excitability of *grey* matter is afforded by the effects of anæsthetics or of interrupted blood-supply; by either of these means cortical excitability is abolished, and that far too rapidly for it to be possible to admit that the subjacent *white* matter has been thus influenced.

The investigation opened up by Fritsch and Hitzig was soon afterwards pursued and extended by Ferrier (1874–78), who employed monkeys, and tested the cortical excitability by weak induced currents. He distinguished between direct and reflex excitability by differences in the character of motor reactions, regarding the reactions obtained by stimulation of the Rolandic area as direct motor reactions, and those obtained by stimulating

the temporal and the occipito-angular areae as reflex reactions, such as might be expected in consequence of auditory or visual sensations. Excitation of the left occipital lobe causes conjugate movement of the two eyes towards the right—*i.e.* ‘the eyes turn, as if an object on the right, casting its image upon the left halves of the retinae, had created an impression in the left occipital lobe.’ In harmony with this view, Schäfer finds that the latent time of the ocular movements is greater to stimulation in the occipital (or visual) area than to stimulation in the frontal (or oculo-motor) area. Ferrier found that direct excitability is limited to the convolutions in the vicinity of the fissure of Rolando, viz. the ascending frontal and ascending parietal. This region he therefore designated as the ‘*motor area*’ of the cortex; and it may be remarked in this place, that Ferrier’s experimental results on monkeys are in general agreement with the deductions that Hughlings Jackson had previously made by

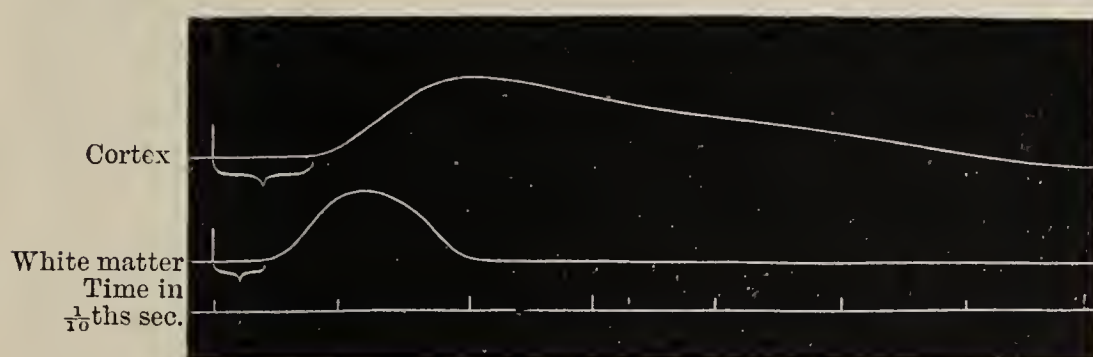


FIG. 293.—MUSCULAR CONTRACTIONS PROVOKED BY CORTICAL AND BY SUB-CORTICAL EXCITATION. (Bubnoff and Heidenhain.)

the clinico-pathological study of epilepsy in man—to wit, that combinations of muscular movements originate in the middle region of the cortex of the brain. Ferrier, moreover, sought to localise the position of sensory centres by observing the sensory defects consequent upon the destruction of different cortical areae; he came to the conclusion that *vision* is represented in and near the angular gyrus, *hearing* in the first temporal convolution, *smell, taste, common and tactile sensation*, in the hippocampal region. But a localisation of sensory regions—assuming that sensory impressions, like motor impulses, are centrally differentiated—is a far more uncertain question than the localisation of motor regions; the margin left to be interpreted—by opinions, or by fancy, or under the influence of preconceived theory—is very large, and consequently many different opinions have been and are held concerning sensory areae. Munk, who subsequently investigated this aspect of the



question (1877), placed vision in the occipital region ; operating on dogs and on monkeys, he found that destruction of the cortex of both occipital lobes caused total blindness, destruction of one occipital lobe caused hemiopia on the same side of the retinae, conjoined with impaired vision of the whole of the opposite eye. These facts are in agreement with the clinical symptoms observed in man : hemiopia sometimes coexists with hemiplegia, and in such case the patient cannot see to his paralysed side. If, for instance, the left hemisphere is at fault, the muscles on the right and the retinae on the left are paralysed, and the right side of the field of vision is blotted out ; moreover, the impairment of vision is greatest for the right eye.

With regard to hearing, Munk was of opinion that the whole of the temporal region is its sensory area ; and he came to this conclusion from an ingenious experiment, in which he avoided the practical difficulties in the way of extirpating both temporal lobes, viz. he destroyed one temporal lobe and the auditory apparatus of the same side ; the dog was thus rendered deaf, and from this result the double inference was drawn—(a) that the temporal lobe is the organ of hearing, (b) that the auditory channels undergo a complete decussation.

Ferrier and Yeo reaffirm the conclusion that destruction of the first temporal convolution on both sides causes complete and permanent loss of hearing, and that entire destruction of the hippocampal region (hippocampus major and gyrus hippocampi) and neighbouring inferior temporal region causes complete anæsthesia of the opposite side of the body. On the other hand, the statement that hearing is localised in the temporal region is formally contradicted by Schäfer, who found no impairment in the hearing of monkeys for weeks after complete destruction of that region on both sides, as verified *post mortem*. We must therefore suspend our opinion with regard to the cortical locus of the sense of hearing.

With regard to *common* and *tactile sensation*, we are in presence of Ferrier's statement, that complete anæsthesia of the opposite side is produced by destruction of the hippocampal region, and of Schäfer and Horsley's statement, that hardly any alteration of sensibility is produced by such lesion, while 'any extensive lesion of the gyrus fornicatus is followed by hemianæsthesia, more or less marked and persistent.'

But the appetite for special centres has lost its edge, and the inference from these statements, to the effect that common and

tactile sensibility are localised in the limbic lobe (gyrus fornicatus et hippocampi), has not met with much acceptance. There are no sufficient grounds for admitting that centres of common sensation are distinct from centres of common motion; the more probable view is that which has been uniformly held by Hitzig and by Munk, to the effect that the Rolandic or 'motor' area is also a sensory area—an arrival-platform of centripetal impulses, and a departure-platform of centrifugal impulses. Munk, indeed, locates 'sensation'—cutaneous, tactile, muscular, tendinous and articular—in the motor area, which he regards as primarily sensory; sensation the cause, motion the effect.

As for *smell*, we can only—in the absence of satisfactory experimental or clinical data—point to the possibility suggested by comparative anatomy: the olfactory lobe and the hippocampal gyrus (uncus) are in anatomical connection, and their greater development has been found to be characteristic of animals with a highly developed sense of smell.

To recapitulate the main results in their order of certainty—

*It is proved that the Rolandic area is 'motor.'*

*It is proved that the occipital area is visual.*

*It is probable that the temporal area is auditory.*

*It is probable that the entire cerebral cortex is 'sensory.'*

**Subdivision of the 'motor' area.**—The Rolandic area has been further subdivided by experimental stimulation into parts which are found to be connected with definite groups of muscular movements of the upper or lower extremities, of the face, neck or trunk. To the localisation thus determined the same qualification applies as that which we recognised in the general question: viz. localisation is not strict, but straggling; there is no exclusive connection between points of cortex and parts of muscle, but a predominant connection between cortical regions and associated or grouped voluntary movements.

On the external aspect of each hemisphere the chief motor loci are as follows:—

Movements of the—

Inferior extremity . . .	Upper portion of Rolandic area
Superior extremity . . .	Middle portion of Rolandic area
Face . . . . .	Lower portion of Rolandic area
Lips and tongue . . . .	Base of third frontal convolution
Head and eyes . . . . .	Base of first and second frontal convolutions



On the mesial aspect of each hemisphere, *i.e.* in the marginal gyrus, the disposition is as follows from before backwards: head, arm, trunk, and leg.

By carefully localised stimulation a still further subdivision of areae is possible, particularly in the case of that of the upper

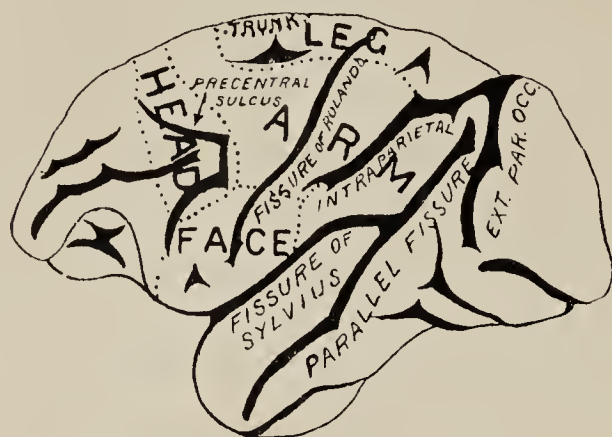


FIG. 294.—LATERAL ASPECT OF MONKEY'S BRAIN, TO ILLUSTRATE CORTICAL LOCALISATION. (Horsley and Schäfer.)

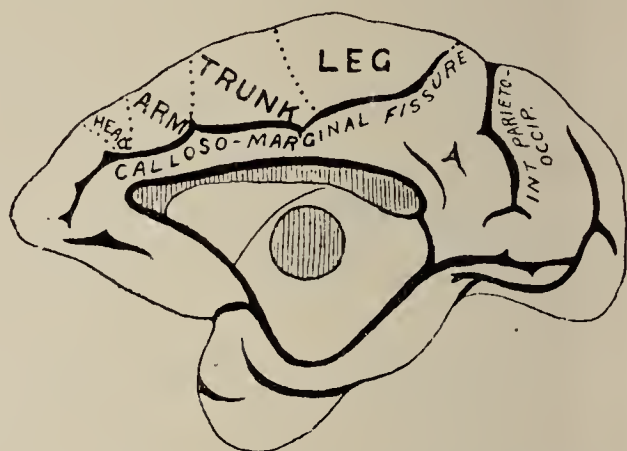


FIG. 295.—MESIAL ASPECT OF MONKEY'S BRAIN, TO ILLUSTRATE CORTICAL LOCALISATION. (Horsley and Schäfer.)

extremity. Particular movements of the arm, forearm, hand, and thumb can be produced by excitation of particular spots 'almost as regularly as definite notes can be sounded on a piano by touching particular keys.'

*Paralytic defects of motion.*—Having recognised these motor reactions as the effects of localised excitation of the cortex, we naturally ask, whether the converse effects will be produced by local destruction of the same parts. We might expect to find a limb completely paralysed after its cortical centre had been destroyed. But this is not the case; the limb can be moved, at first sight its movements seem perfect, it is certainly not completely paralysed. On closer examination, however, some defect of movement will probably be detected, some clumsiness of manœuvre indicating that certain movements are inaccurately co-ordinated, and these imperfect movements will probably be such as are normally most highly specialised, and *not* such as are most profoundly automatic. The limb is not paralysed, but its movements are not perfect, and it is not skilfully used. But even this amount of defect is not apt to be permanent; the apparent awkwardness soon wears off, movements seem to be re-learned; the clinical history of a dog or of a monkey having suffered a removal of some portion of the motor area altogether negatives a strict localisation of function, but forcibly suggests its local concentration; and comparing the effects on animals high and low in the scale, we recognise that that concen-

tration has reached its highest expression in the human brain ; in a given animal, the larger the lesion the more permanent is the defect ; but in different animals a given lesion produces a more permanent defect in the animal highest in the scale ; in man, the least extensive lesion produces the most permanent defect, and the defect produced by a considerable loss of cortex is apt to persist throughout life.

**Occipito-retinal correspondence.**—According to Munk, different parts of the retina are in functional connection with different

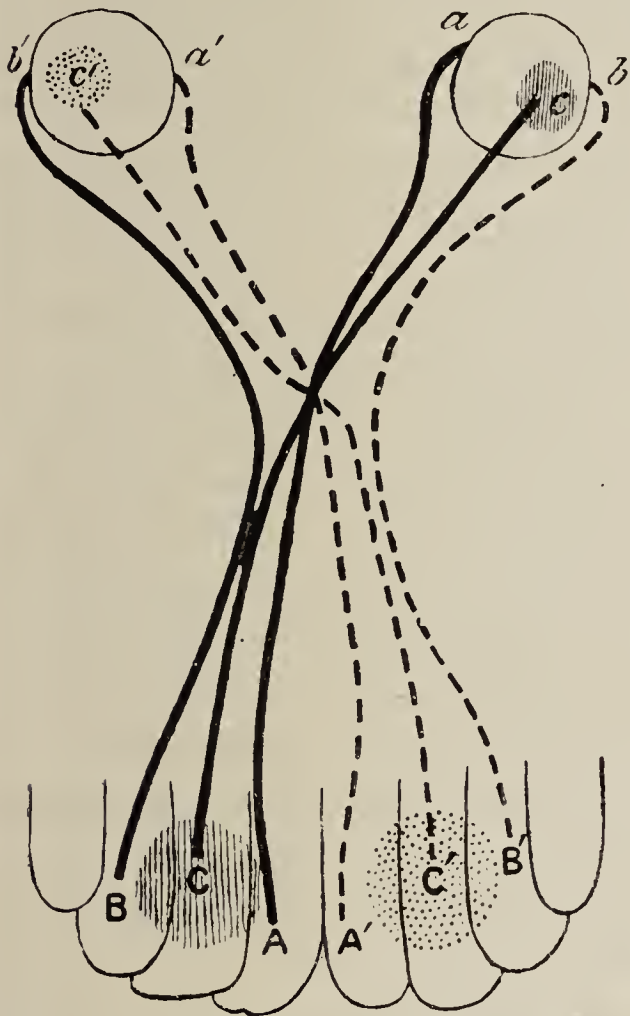


FIG. 296.—OCCIPITO-RETINAL CONNECTIONS, ACCORDING TO MUNK, FROM EXPERIMENTS ON DOGS AND MONKEYS.

Each occipital lobe, A, B, C, is connected with *a* and *c* of the opposite retina, and with *b* of the same side.

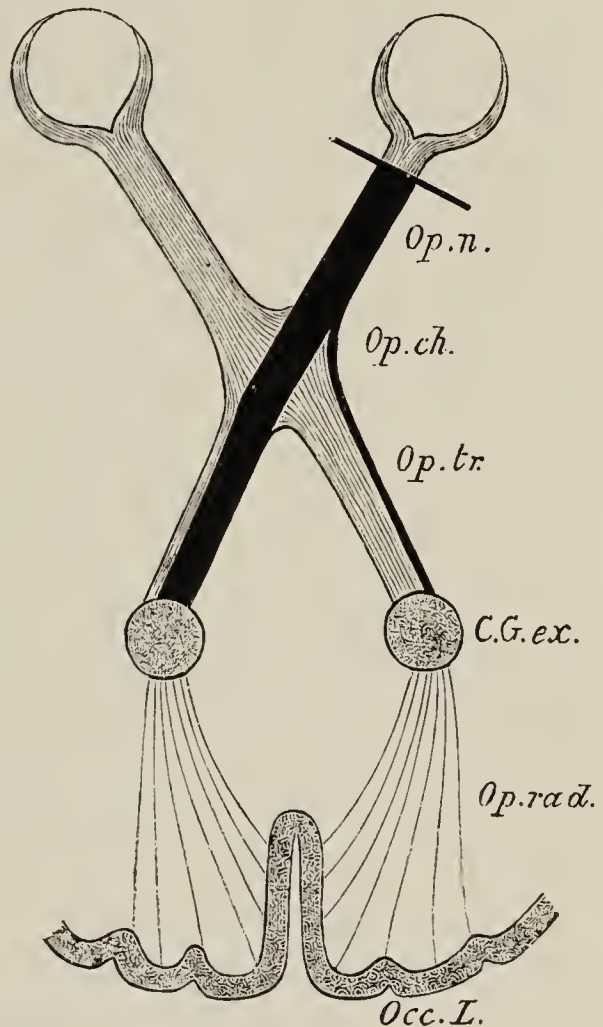


FIG. 297.—DEGENERATION OF OPTIC NERVE AND TRACTS CONSEQUENT UPON SECTION OF THE NERVE (OR ENUCLEATION OF THE EYEBALL).

The degeneration includes a greater part of the opposite tract and a smaller part of the tract on the same side.

parts of the visual area of the cerebral cortex, the retina being, as it were, 'projected' upon the occipital region of the opposite side, so that the upper retinal segment corresponds with the antero-occipital area, the lower retinal segment with the postero-occipital area, the central spot of the retina with about the centre of the occipital area. Schäfer, while supporting the principle of retino-cortical projection, considers that the visual areae of the two sides overlap in such a way that the central spots of both retinae are projected together near the middle line. A de-



termination of the field of vision on animals is obviously a matter of extreme uncertainty; and clinico-pathological records on man do not justify any detailed mapping-out of retino-occipital correspondence. Even the principal statement that each occipital lobe represents the opposite retina, and with it the doctrine—clinical as well as experimental—that destruction of the occipital cortex on one side entails blindness of the opposite eye, are now discredited and all but supplanted by the more probable view that in man, dog, and monkey alike, the left sides of the two retinae are represented in the left occipital cortex, the right sides in the right. This view is presented in Munk's later diagram (fig. 297) exhibiting as the main effect of destruction of the occipital lobe on one side, paralysis of the two retinae on that side, most extensive in the opposite eye (homonymous bilateral hemianopsia). It is moreover in harmony with our knowledge of optic tract degeneration consequent upon destruction of the eyeball or division of the optic nerve; in this connection the principal facts (first established by Waller in 1856, confirmed and extended by Gudden in 1874) are that the retina is the trophic centre of optic fibres, and that the latter when separated from their trophic centre undergo degeneration extending along a large part of the opposite optic tract, and a smaller part of the tract on the same side up to the basal ganglia. From clinico-pathological sources the chief significant points are that in cases of homonymous bilateral hemianopsia, cortical lesions have most frequently been found in the vicinity of the calcarine fissure, *e.g.* in the cuneate lobule (Henschen), and that in such cases the blind area has been found to be more extensive of the opposite eye, but not to involve central vision of either eye. This last point signifies that the fibres from the central spot of distinct vision, undergo semi-decussation, and that each macula is in functional connection with both sides of the cerebral cortex. All the points summarised above are in conformity with the partial decussation at the chiasma, and we may be disposed to admit as a natural relation that the two sides of the retinae upon which lateral images are cast should be chiefly connected with that side of the brain whence chiefly proceed the muscular reactions by which objects are repelled or seized; an assailant from the left, imaged on the right halves of the retinae, is 'seen' by virtue of the right calcarine cortex, and repelled it may be by an instinctive muscular reaction of the left side aroused from the right Rolandic

cortex. As regards the diagnostic localisation of lesions, it may be remarked that complete blindness of one eye indicates lesion below the chiasma, homonymous half-blindness of both eyes lesion above the chiasma (optic tract, external geniculate body, calcarine cortex), whereas asymmetrical affections of one or both eyes are probably due to partial lesions anywhere on the retino-cortical path, excepting only the cortex itself.

**Epilepsy.**—The minute study of epileptic spasms has contributed largely to our knowledge of cerebral mechanism. Clinically, the form of epilepsy which most nearly concerns the physiologist is that known as Jacksonian, or cortical epilepsy, which has a closely similar counterpart in the epileptoid attacks produced on animals by experimental stimulation of the cortex. A typical epileptic fit of cortical origin is divisible into the definite stages (1) of the warning sensation, or *aura*; (2) the fit proper, or period of excessive muscular discharge; and (3) a post-epileptic stage, characterised by muscular debility, which may amount to actual paralysis, and which is referable to an exhaustion of ‘discharged’ cortex. The entire phenomenon is comparable to a protracted series of explosions, arising in an over-irritable or definitely irritated spot of the cortex as its focus of origin. The character of the warning and the mode of commencement of a fit give a clue to the actual seat of irritation; the order of invasion and the character of the convulsion are the outward sign of the directions along which the original cortical disturbance has spread. The warning sensation may consist in a peculiar sensation in the muscles of the limbs in which the convulsion is about to commence, or it may take all kinds of subjective forms, from a simple flash of light, or a sound, smell, or taste, to an elaborate representation of scenery or of music. Such muscular, visual, or auditory auræ, if well characterised, are to be taken as symptoms pointing to the area within which irritability is overstepping the boundary-line into actual irritation. The fit proper—clinically divided into a short tonic period, during which muscular contraction is at its height, and a longer clonic period, during which muscular contraction is resolving itself—is significant of excessive discharge from the Rolandic area of the cortex. Physiologically, the feature of greatest interest is the order in which the limbs become affected at the very commencement of the fit. It may commence in any one limb. Ordinarily it does so by twitchings of the highly specialised extremities most used in voluntary movement, viz. the thumb and fingers, and thereafter



involves the forearm and arm, the opposite upper extremity, the face, and finally the lower extremities. But the order of invasion is by no means invariable. The part first invaded does, however, furnish valuable hints towards determining the precise seat for operative interference in cases where there may be reasonable hope of cure by removal of irritation: a fit beginning in the left great toe points to a discharging lesion of the upper part of the right Rolandic area; a fit beginning in the right hand points to a discharging lesion about the middle of that area.

The variability of the order of invasion deserves attention. We are naturally led to inquire whether the diffusion occurs in the cortex, or in subordinate bulbo-spinal centres. The answer to this question is, that the diffusion probably occurs in the cortex *and* in subordinate centres. Or there may be no diffusion, the epileptoid spasms remaining limited to a single limb, or to one side of the body, as a monospasm or as a hemispasm.

The phenomena of experimental epilepsy confirm this interpretation, and fix it in certain particulars. An epileptoid attack remaining locally restricted, or taking place as the prelude to a general convulsion of the body, is apt to occur whenever experimental stimulation of the Rolandic area is at all prolonged or strong; the attack occurs or begins in the limb or part functionally connected with the particular cortical area excited; and if it become generalised, the order of invasion is, as in clinical cases, a variable one. Its more usual but by no means regular order is—supposing the spasm to begin in the right upper extremity—as follows: right upper extremity + right lower extremity + left upper and lower extremities, *i.e.* the invasion is longitudinal before it is transverse. This order favours the view that the diffusion has occurred in the cortex of one side rather than in the spinal cord, where, as we have seen, diffusion of reflex stimuli is transverse before it is longitudinal (p. 497).

The great part played by the cortex in the extension of an epileptoid storm is, moreover, demonstrated in the experiments of Munk and of Heidenhain. Munk arrested the partial epilepsy which he had excited in a single limb, by rapidly excising the cortical seat of commotion. Heidenhain went further, and in an early stage of a general epileptic attack put a single limb to rest by excising the cortical area by which it was governed. According to all experimenters, no epileptoid attack can be produced after destruction of the cortex of both hemispheres.

But the doctrine that epileptoid diffusion is *exclusively* cortical

is not justified by further experiments. (1) Epileptoid convulsions may extend to parts of the body after removal of the cortex that is in relation with them. In this case the storm provoked in uninjured cortex has involved subordinate centres, and has there become diffused. (2) Epileptoid convulsions may be provoked by stimulation of white matter after removal of the overlying cortex. In this case excitation of white matter has reached uninjured grey matter through commissural fibres, and there provoked a storm which has become diffused in subordinate centres. These points will be appreciated by following the lines of the diagram.

**Aphasia**—literally loss of speech, more precisely loss of language—is one of the consequences of cerebral disease, and the discovery that such disease usually involves the third frontal convolution of the left hemisphere, was the first definite step in the localisation of cerebral diseases and of cerebral functions. The minute analysis of the various forms and symptoms of aphasia is thus an important source of information in the study of cerebral mechanism.

The subject of aphasia in its most typical form is unable to express ideas correctly in words, either spoken or written, and this inability may be of any degree, from a transient or occasional misuse of a word, such as most people have experienced after hard mental work, to an habitual misuse of words, or to their complete loss. The names of persons and of concrete things are the first to go; the verbal symbols of complex and abstract ideas cling closer and longer. Such an affection is termed *motor aphasia*, and its subject, though possessing ideas, cannot emit them for want of words, or emits them in wrong words; the ideas have no clothes to go out in, or they go out in the wrong clothes.<sup>1</sup> It is

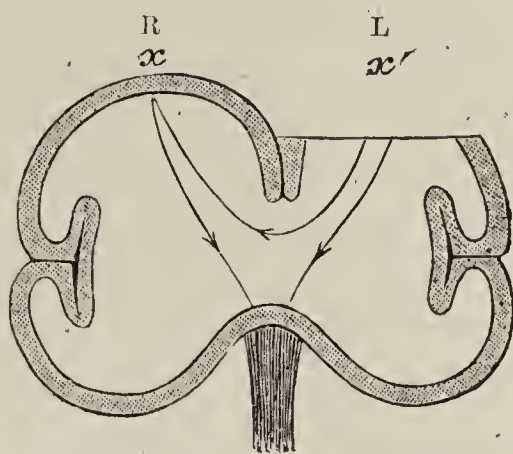


FIG. 298.—EPILEPTOID DIFFUSION.

The motor cortex of the left hemisphere is removed. (1) Stimulation  $x$  of the motor cortex on the opposite side has provoked an epileptoid attack; diffusion must have taken place in the bulbo-spinal grey matter. (2) Stimulation  $x'$  of the exposed white matter from the motor cortex of the left hemisphere has reached the right hemisphere through the corpus callosum, and has provoked an epileptoid attack, which has arisen in the right motor cortex, and must have diffused in the bulbo-spinal grey matter. (3) No epileptoid attack could be produced by stimulation of white matter after removal of the motor cortex on both sides.

<sup>1</sup> A word is a complex bundle of movements—of the tongue and lips in *speech*, of the hands in *writing*, of various limbs in the *pantomime* sometimes used to



this form of aphasia which has been found associated with disease of the left third frontal convolution.

The term aphasia is also applied to a form of the disease in which the patient can express ideas in monologue, but cannot answer questions, this inability being due to the non-recognition of the words that he hears spoken or sees written. The defect may vary in degree from a transient absent-minded pause, such as occurs in the experience of most people, to the complete blank non-recognition of words. Such an affection is termed *sensory aphasia*, and is often spoken of as *word-blindness* or *word-deafness*, according as its subject fails to recognise words by sight or by sound. This form of aphasia has been found of late years associated with disease of the visual or auditory regions of the cortex, *i.e.* of the occipital or of the temporal regions. The condition of a man, the subject of sensory aphasia, is strictly comparable with that of a dog from which the cerebral cortex has been removed: the aphasic man can see objects and hear sounds, but cannot recognise the meaning of written or of spoken words; the dog likewise can see and hear, but does not recognise the idea of a bone placed before him, nor the idea of the crack of a whip; he exhibits neither desire of the one nor fear of the other.

Let us follow the steps by which a rational answer is given to a question. (1) The subject must hear or see a word or words. (2) The word heard or seen must excite an appropriate sensory idea. (3) The sensory idea must give rise to a consequent motor idea, *i.e.* a word or words. (4) The word must be spoken.

No answer will be given if the subject be deaf or blind (1), or dumb (4); he is not on that account said to be aphasic. But he is so if he has no word with which to dress his outgoing idea (motor aphasia). Destruction of the third left frontal convolution, where his ideas are dressed, or destruction of the motor fibres from this convolution, causes such wordlessness. The destruction may be actual or functional, complete or incomplete; if incomplete, the symptom will be incomplete; ideas few or many will be sent out dressed in wrong words. Again, he is aphasic

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denote mental ideas and to form propositions. In its restricted sense language is speech by words spoken, written, or gesticulated; in an extended sense language is any and all muscular action expressive and indicative of states of mind. A falsehood is any word or action voluntarily suggestive of erroneous inference on the part of listener or observer.

if he cannot recognise the words in which another man's ideas are expressed (sensory aphasia), and theoretically he will also be aphasic if communicating channels be interrupted between the sensory, visual, and auditory regions and the motor, or word region proper.

The study of aphasia further informs us in certain other important particulars. Aphasia is commonly associated with *right hemiplegia*, the common cause being obstruction of the left Sylvian artery. Aphasia due to this cause is generally of a mixed character, compounded of motor and of sensory symptoms, *i.e.* defects of emission (motor aphasia) and defects of recognition (sensory aphasia). The right Sylvian artery is seldom blocked, but when it is, left hemiplegia is its consequence, and aphasia is absent, except in rare cases, and it is said that most of these rare cases have occurred in left-handed people. These several facts go to show that the left side of the brain is normally the leading side, not only as regards motion on the right side, but also, and more so, as regards speech; but that, as an exception, this precedence may be reversed, the right side of the brain being the leader, giving superiority to the movements on the left side, and taking charge of speech.

The cortical centre of the closely associated function of vocalisation coincides with that of verbalisation, but in this case the centre is apparently bilateral—excitation on one side of the brain giving vocal movements on both sides, and destruction of both frontal lobes being necessary before these movements are abolished.

*Recovery.*—Aphasia may be permanent or may gradually fade away; a patient, speechless during the first few days after seizure, slowly recovers the use of proper language, and if the lesion has not been too profound, his recovery may to all appearance be complete. To explain this fact. In most cases it is doubtless due to restored circulation and nutrition. But in others we are tempted to suppose that the right side of the brain has gradually taken up the duty that normally belonged to the left side. The sensori-motor apparatus concerned in the perception and expression of language holds on the left side, as compared with the right, a functional rank which is analogous with that held by captain and lieutenant; normally, the latter acts in agreement with and upon the initiative of the former, but may on emergency succeed to and carry out the initiative action that normally belonged to the leader. In other cases, again,



*i.e.* when aphasia is attributable to destruction of communicating channels between the left frontal gyrus and subordinate bulbar nuclei, it is possible that the rapid recovery that may occur is owing to the left side resuming action through the commissural fibres which form its communication with the opposite side of the brain. But these are surmises that cannot be brought to trial.

The acquisition of language and its loss by *unilateral* cortical lesion illustrate a principle of far-reaching significance. Language is in the highest degree a special function, evolving last in the animal scale—taking form comparatively late in the individual life—most variable with varying education—first to show signs of failure in the degradation of cerebral function. The general functions and their organs are formed earlier than the special functions and their organs; the special functions and their organs are the first to fail in the dissolution of the formed organism. To use Hughlings Jackson's expression, language is a highest-level function; it comes last and goes first.

*Deaf-mutes.*—The modern or 'oral' education of deaf-mutes illustrates the acquisition of language under difficulties. In such people congenital deafness is the primary defect; muteness is its consequence, because the speech mechanism, although anatomically perfect, is deprived of education by sound, and remains undeveloped. In the place of articulate speech a deaf-mute employs a code of signals or intellectual pantomime for the exchange of ideas. Until within recent years this imperfect form of language was the only one taught; but it has been found that deaf-mutes can learn to recognise ordinary language, which they cannot hear, by seeing the lip-movements of a person who is distinctly articulating; and, further, that by close imitation of these lip-movements they may learn to articulate distinctly in ordinary language. Such people learn speech by sight, whereas normal persons learn speech by sound, and the education of a person born deaf may be carried to such a pitch of perfection that the defect may pass unnoticed in ordinary conversation.

Puppies, or even dogs, are rendered dumb by destruction of the internal ear; a dog learns to bark by hearing other dogs bark; failing sound, he does not apparently learn to bark by sight, and unlike a man in this respect (who does not forget how to speak when he has become deaf), he may even forget how to bark if he loses his hearing late in life (Danilewski).

In the foregoing account, and particularly in the short statements on p. 541, the terms motor and sensory are used in their ordinary sense, but 'without prejudice,' and without implication of any theories with regard to the essential nature of the cortical processes so designated. It should, however, be added that opinions and interpretations of the intimate nature of the process take very various shapes, and that inferences from facts are clothed in many kinds of verbal garments. Hitzig and Bastian interpret the results of excitation of the motor area as evidence that it is the central organ of muscular sense, and the latter maintains that impairment of 'muscular sense' is associated with the impairment of motility caused by lesion of the Rolandic area. Flechsig, from entirely independent data, comes to the conclusion that the motor cortex is sensori-motor. Golgi on histological grounds, and Münsterberg on philosophical principles, come to the same conclusion. Munk sees in his own results evidence that it is a sensori-motor area (*Fühlsphäre*); he finds that not only movements, but also sensations from the skin and from the muscles, are interfered with when portions of the motor cortex are extirpated, and he regards the defect of movement as secondary to the defect of sensation. With regard to the occipital area, he distinguishes a central portion, c, extirpation of which gives psychical blindness (*Seelenblindheit*), i.e. non-recognition of seen objects, from the surrounding portions, A B, extirpation of which (added to that of c) gives total blindness (*Rindenblindheit*). In the temporal area he distinguishes a central portion, giving psychical deafness (*Seelentaubheit*), from a surrounding portion, giving total deafness (*Rindentaubheit*).

The 'sensori-motor' conclusion is, indeed, an inevitable one; any motor or discharging centre must also be a sensory or receiving centre; it must be excited as well as excite. Any 'sensory' centre must also be motor, directly or indirectly; else we could have no objective tokens of sensation; every centre, whether called motor or sensory, is *terminus ad quem* as well as *terminus a quo*. 'Sense sure you have, else could you not have motion.' Hughlings Jackson from the clinical standpoint particularly insists upon the 'sensori-motor' character of all centres, as opposed to the crude conception of 'motor' centres, and in his hierarchical schema of central nervous mechanism represents sensori-motor centres in three grades—(1) highest-level centres (the præfrontal cerebral cortex), (2) middle-level centres (the Rolandic cerebral cortex), (3) lowest-level centres (the pons, bulb, and cord). This classification is practically the same as that which we have followed in the division into cerebral and spinal, with the additional subdivision of cerebral into upper and lower; and although, pathologically, the division may be legitimised by the analysis of epileptic and other phenomena, experimental physiology does not yet embrace these facts, nor recognise a distinction between supreme and subordinate cortex cerebri. But the actual localisation of three levels is quite



secondary to their ideal conception; the contemplation of human conduct, normal and abnormal, fully warrants that conception, and it is of secondary importance whether the first two levels are both cortical, or whether one is cortical and the other sub-cortical.

Munk's conception is essentially similar to Jackson's, with the addition of an attempt at an actual localisation of two cerebral levels; but the terms he has used have obstructive connotations, and we should accept them more readily if they were more vague. If for *Seelenblindheit* we substitute 'high-level blindness,' for *Rindenblindheit* 'high + middle-level blindness,' we seem to understand better what is meant, and what is the drift of the experiments in point. And in considering the centripetal aspect of nervous processes, we may with advantage appropriate to the three-level scheme the three terms impression, sensation, perception—using 'impression' as the lowest-level term, 'sensation' as the middle-level term, 'perception' as the highest-level term; and taking impression (*Eindruck*) to denote an effect that does not reach consciousness, sensation (*Empfindung*) to denote a felt impression, perception (*Wahrnehmung*) to denote a sensation in relation with its felt circumstances, *i.e.* the group of associated sensations that complete the mental picture or representation of the moment (*Vorstellung*), forming what may be designated as a field of attention.

The reaction time is the interval between the application of a stimulus and the responsive signal indicating that the stimulus has been 'felt.' This interval is conveniently measured by arranging an electro-magnet to mark on a revolving cylinder (1) the moment when a tactile, auditory, or visual stimulus is applied, and (2) the signalling movement by which the person experimented upon indicates that he has felt the stimulus. The reaction time varies in different subjects, with different modes of stimulation, and with different degrees of attention and of health, between ten and twenty hundredths of a second.

The total reaction time is composed of (1) the time occupied in conduction up sensory and down motor nerves, and (2) the time occupied in the central elaboration, during which entering impression gives rise to outgoing impulse. Thus, in the simplest case, where the skin of the hand is stimulated by an induction shock, and the signal given by the same or by the opposite hand, the interval (say  $\cdot 15''$ ) between stimulus and signal is made up of the time of conduction along sensory nerve (say  $\cdot 02''$ ) + the time of conduction along motor nerve and the muscular latency (say  $\cdot 03''$ ) + the time of cerebral elaboration, *i.e.* perception of sensation and formation of volition (say  $\cdot 10''$ ). It is the last of these

three factors that varies most—with idiosyncrasy, attention, health, &c.—and it is partly on this account, partly because the sensibility differs at different parts of the skin, that measurements of the reaction time with cutaneous stimulation near and far from the head give no admissible data for estimating rate of conduction along sensory nerves. The reaction time is shorter when attention is concentrated upon the intended movement than when it is concentrated upon the expected stimulus; this signifies that the formation of volition involves more labour than the perception of sensation, the preparatory influence of attention being more effective if bestowed upon the outgoing than upon the incoming event.

Average values of the reaction time are :

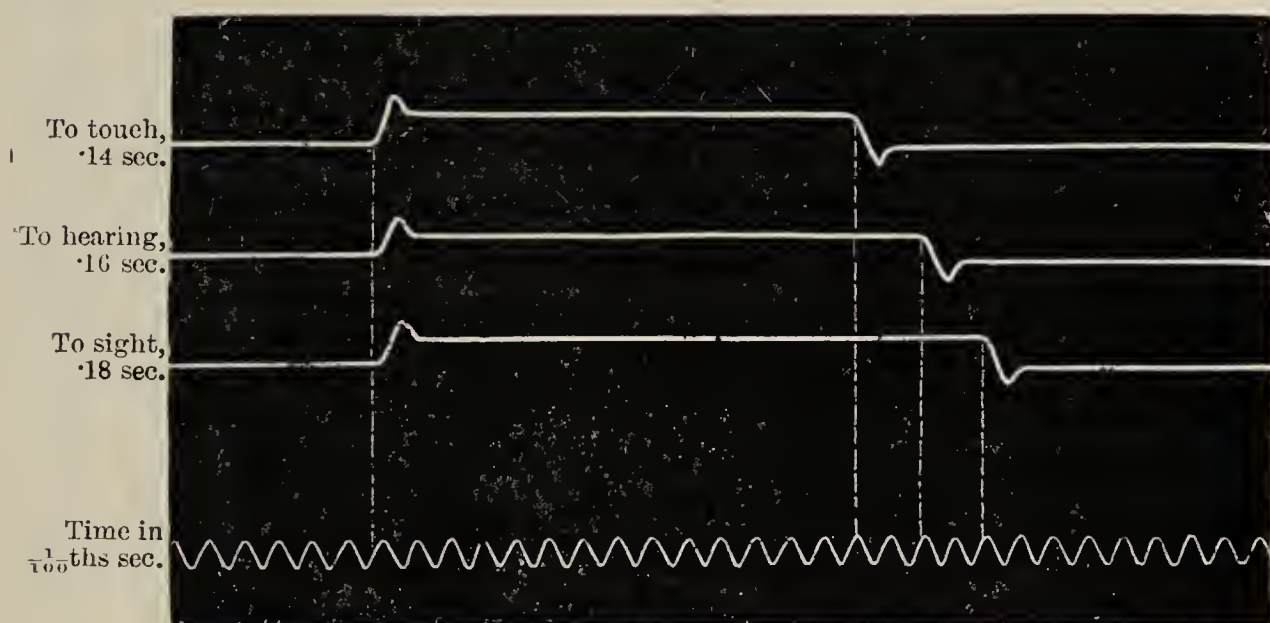


FIG. 299.

With regard to individuality, it is not the case that persons reputed quick and 'wide-awake' have a shorter reaction time than persons of an apparently sleepy and phlegmatic temperament. 'Quick' people often have a long reaction time; 'slow' people often have a short reaction time; the relation, although common, is, however, not constant enough to constitute the rule. It is said that women are, on the average, quicker than men, and negroes quicker than whites. Experiments can easily be devised so as to yield an approximate estimate of the shortest time required to discriminate between two sensations (discrimination time), and of the shortest time required to determine an act of volition (volition time). For instance, the hand of a person, with bandaged eyes, on whom the simple reaction time to touch has been determined to be, say,  $\cdot 15''$ , is stimulated on the little finger or on the thumb, with the understanding that



only one of these stimuli is to be signalled ; the reaction time is now found to be, say,  $\cdot 17''$ , from which the conclusion is drawn that  $\cdot 02''$  was the discrimination time, *i.e.* that required to distinguish between the two different sensations. Or the experiment is conducted with two signals, on the understanding that one signal is to be used when the little finger is touched, and the other when the thumb is touched ; the reaction time under these conditions being found to be  $\cdot 20''$ , is considered to be the sum of  $\cdot 15''$ , the simple reaction time,  $+ \cdot 02''$ , the discrimination time,  $+ \cdot 03''$ , the volition time. The experiments may be still further complicated in a variety of ways, to measure the time of recognition of letters, words, simple ideas, &c., and are then distinguished by the title 'psychometric.'

*Stimulation and sensation. The Weber-Fechner law.*—Our measure of sensation being necessarily subjective, consists at best in the perception of the least observable difference between two sensations.<sup>1</sup> We assume that this least observable difference or differential sensation is a constant sensory quantity, or one sensation unit ; *e.g.* the differential sensation between two weights of 170 and 180 grammes, and the differential sensation between two weights of 1,700 and 1,800 grammes are found by experiment to be 'least observable differences'—that is to say, equal sensations. Each such sensation is then assumed to be one unit, and, in accordance with Fechner's exposition, the assumption includes the differential sensation between zero and the smallest appreciable quantity of any stimulus—this minimum perceptible sensation being, in fact, the smallest observable difference between zero and something. This assumption is the basis of psychophysics. Within certain limits, the strength of sensation increases with the strength of stimulation, and the relation is such that each equal increment of sensation requires an increasing increment of stimulation ; *e.g.* a given sound of intensity = 1, producing a sensation = 1, must be increased to an intensity =  $\frac{4}{3}$  before any difference becomes perceptible, to an intensity =  $\frac{4}{3} \times \frac{4}{3}$  or  $(\frac{4}{3})^2$  before a second difference is perceptible, to an intensity =  $\frac{4}{3} \times \frac{4}{3} \times \frac{4}{3}$  or  $(\frac{4}{3})^3$  for

<sup>1</sup> Nothing can be perceived otherwise than by comparison with something else ; the most elementary sensation is the resultant of a *ratio* between *two* sensificatory terms, *a* and *b*, and not that of a single term, *a* or *b*. Something is perceived in comparison with nothing or with something else, and the resultant of the ratio in consciousness is a sensation. The highest forms of sensation are equally reducible to a ratio between *two* terms, each of which may, however, be a highly compound ratio.

a third difference, to an intensity  $= \frac{4}{3} \times \frac{4}{3} \times \frac{4}{3} \times \frac{4}{3}$  or  $(\frac{4}{3})^4$  for a fourth difference, &c.<sup>1</sup> Thus the relative magnitudes of stimulus and of sensation will be expressed by the following numbers, or graphically by fig. 300 :

Sensation $y$ .	1	2	3	4	5	6	7	8	9	10	&c. $n$
Stimulus $x$ .	1	$\frac{4}{3}$	$(\frac{4}{3})^2$	$(\frac{4}{3})^3$	$(\frac{4}{3})^4$	$(\frac{4}{3})^5$	$(\frac{4}{3})^6$	$(\frac{4}{3})^7$	$(\frac{4}{3})^8$	$(\frac{4}{3})^9$	&c. $(\frac{4}{3})^{n-1}$
$y$ or	1	1.33	1.77	2.37	3.16	4.21	5.62	7.5	10	13.3	&c.

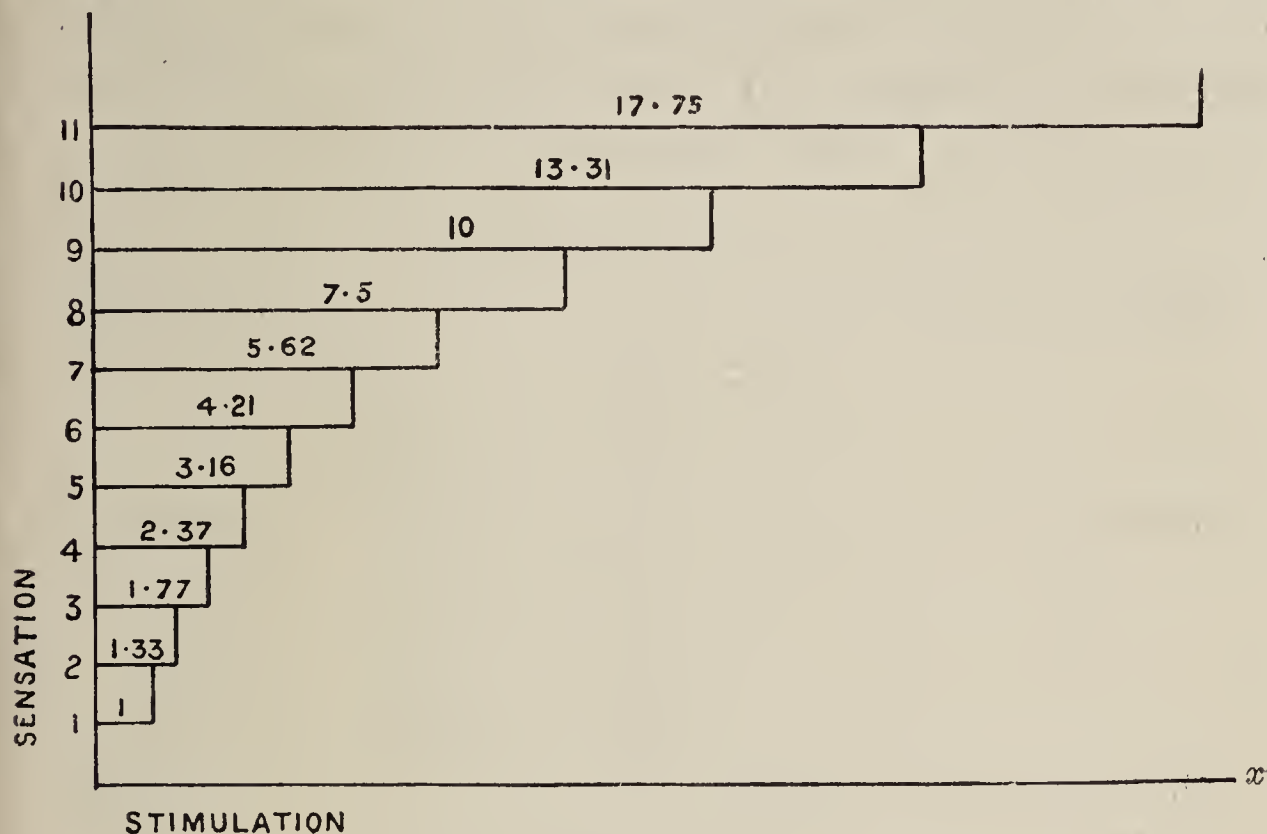


FIG. 300.

Thus we may conceive that any sensation mounts from zero to a maximum by a series of equal units, being the least perceptible differences between its successive magnitudes, beginning from the first magnitude, or minimum perceptible difference between nothing and something. In fig. 300, where sensory units are represented along the ordinate and stimulation units along the abscissa, we see the progressive increase of the stimulation increments required for each successive step of sensation increase. Otherwise expressed, and without the assumption that sensation units may be treated as of constant magnitude, the relation may be put thus: Within a certain range the smallest perceptible difference of stimulation is a constant fraction of the original stimulation. This fraction varies

<sup>1</sup> Sensation increases arithmetically, while stimulation increases geometrically; the equation to the curve, expressing their ratio, is  $x = a^y$ ; i.e. the curve is logarithmic, and the sensation  $y$  varies as the logarithm of the stimulus  $x$ , since  $y = \log_a x$ ; this is Weber's law. Fechner's formula is  $y = k (\log x - \log x_0)$ , where  $y$  is the sensation,  $k$  an experimental constant (differing with the nature of the sense-organ),  $x$  the stimulus, and  $x_0$  the liminal intensity (*reizschwelle*).



with different sense-organs ; thus, for *light* the smallest perceptible difference is  $\frac{1}{100}$ , for *weight* it is  $\frac{1}{17}$ , for *sound* it is  $\frac{1}{3}$ . This relation, known as Weber's law, is a phenomenon of everyday experience, affecting every province of human feeling, setting a limit to pains and pleasures, yet permitting the appreciation of differences within a very extensive range of stimulation intensity. Referring to the curve given above, we see how rapidly a maximum of sensation is approached with increasing stimulation intensity, and a glance at the table below shows us within what a wide range of light intensity we habitually live.

	Sensation units	Stimulation units	
Minimum perceptible . . . . .	1	1	$\frac{1}{4000}$ candle
	11	1.1105	
	101	2.704	
	201	7.314	
	301	19.789	
Moonlight . . . . .	370	40	$\frac{1}{100}$ candle
	401	53	
	501	144	
	601	390	
	701	1059	
	801	2865	
White paper lighted by candle at 1 foot	834	4000	1 candle
	901	7749	
	1001	20960	
	1101	56695	
	1157	100000	25 candles
	1201	153350	
	1301	414790	
	1388	1000000	250 candles
	1401	1121900	
	1501	3027900	
White paper lighted by sun . . .	1601	8208100	
Full sunlight . . . . .	1701	22201000	5,555 candles

*Weight-discrimination.*—In the distinction between lighter and heavier weights, it is generally assumed that the physiological sensation caused by ‘lifting’ weights is due exclusively to peripheral strain and pressure—of the muscles, of the tendons, of the articulations, and of the skin. Whatever be the view taken of the actual constitution of the sensation of movement or effort, we can make observations as to its delicacy, and in so doing acquaint ourselves with the three chief methods of investigation that it is possible to follow. We may, as in the case of light, employ the

method of (1) *minimum perceptible difference*, and find the smallest difference that we can appreciate between two weights ; but this will be found to yield very variable results, for we must either make the comparison simultaneously with two hands, or successively with one hand. Or we may proceed as follows : we may first lift a given weight, and then lift a second variable weight, which we adjust until it is judged to be equal to the first ; there will probably be an error in the estimate ; the sum of several errors is taken, and divided by the number of trials ; this is (2) the method of *mean error*, and although, with the adoption of certain corrections, it is superior in accuracy to the first method, it does not lend itself to mathematical analysis. The best method is (3) that of *right and wrong cases* ; in principle it is an application of the probability calculus to large numbers of trials ; in practice it is as follows : two weights differing by a certain fraction are given to be distinguished ; a large number of trials are made with the same pair ; if the difference is within certain limits, a certain number of right and of wrong answers will be given ; the greater the difference, the greater the proportion of right answers to the total number of trials. With sufficiently large numbers of trials (say 1,000 for each pair of weights), a measure of sensibility may be calculated from this proportion.<sup>1</sup>

Having learned that the relation between stimulus and sensation is such that equal increments of the former cause diminishing increments of the latter, we are naturally led to ask whether the transforming factor occurs in physiological or in psychological territory. Is the material sensificatory brain-change proportional to the external stimulus, and the sensation some geometric function of that change, *or* is the material change itself a geometric function of the stimulus, and attended with sensation in direct proportion to its own magnitude ? Fechner and Wundt subscribe the first alternative, to the effect that the transforming factor is psychological, *i.e.* between the sensificatory change and the sensation. Pfeffer, Delbœuf, and others, hold that the disproportion is physiological, *i.e.* between the stimulus and the sensificatory change. The fact that a similar disproportion between cause and effect is witnessed in the purely

<sup>1</sup> The necessary calculation is reduced to a simple operation of arithmetic by making use of Fechner's Fundamental Table, which gives the numerical values of 'the measure of exactness,' *i.e.* 'the measure of sensibility for different values of  $r : n$ ' (*Psychophysik*. I. p. 108).



physiological instances of muscle (p. 328) and of the retina (p. 401) weighs in favour of this second alternative.

**Muscular sense, sense of movement, sense of effort,** are terms used to denote the sensation that accompanies all voluntary muscular contractions. The factors of this composite sensation are —(1) *peripheral*, arising from variations of tension and of pressure in the moved organs, *i.e.* in tendons, ligaments, cartilages, possibly in the muscle itself, certainly in the skin of the part; (2) *central*, arising from material changes accompanying the elaboration and emission of motor impulses.

The view that the central state contributes to the sensation is supported (1) by the fact that weight-discrimination is far more delicate by voluntary than by artificially excited contraction, (2) by the consideration of fatigue changes, which are demonstrably central as well as peripheral. Admitting that material changes *after action* occur in the same parts as material changes *with action*, we may learn the incidence of the effect by studying the incidence of the after-effect. Now the after-effect of muscular action, subjectively felt as fatigue, is demonstrable by objective signs at the centre as well as at the periphery; from which we infer that the motor effect, which is subjectively felt as the sensation of movement or of effort, is associated with material changes at the centre as well as at the periphery. This is to say, that the sensation called ‘muscular sense,’ ‘sense of movement,’ ‘sense of effort,’ is associated with changes *at the centre and at the periphery*; we have no right to say that it is caused by changes in either of these situations to the exclusion of the other (w.)

**Sensory judgments, illusions.**—Comparisons and judgments, unconscious as well as conscious, are performed in the brain, and consist in the collation and comparison of sensations, which may be simultaneous or successive. We shall refer, in illustration of this subject, almost exclusively to the sense of vision; it is the most important and most cerebral of the senses, and the most accessible to experimental analysis; but it should be remembered that channels of sensation other than visual contribute sensations, which are compounded in consciousness, and which supplement or correct visual and other judgments.

The popular expressions, ‘a correct eye,’ ‘sense of proportion,’ are indicative of composite central qualities, partly inherited, partly acquired by education through the senses.

It is not sufficient to possess sensory instruments: we must also be able to interpret their indications by cerebral inference. The simplest case to take in illustration of cerebral inference is the way in which we judge of the size of objects, and of their *distance* from the eye. The size we attribute to an object is the resultant of two factors—the actual size of its retinal image, the known or apparent distance from the eye. The distance from the eye at which we estimate that an object is situated, is likewise the resultant of two factors—the actual size of its retinal image, the known or presumed size of the object. When we have a small retinal image of an object we know to be large, we judge it to be far off; when we have a large retinal image of an object we know to be small, we judge it to be close to the eye. An after-image appears larger or smaller according as the gaze is turned to a more or less distant surface, the branded area of the retina corresponding to a larger or smaller area of the more or less distant surface.

The standard of reference that we unconsciously or consciously employ in forming a conclusion from distance to size, or from size to distance, is the cerebral resultant of previously felt relations between size and distance—the personal ratio drawn from our stock of similar experience. Formally expressed, the major or general premise is our cerebral ratio of previous similar *felt size: felt distance*; the minor premise is one or other of these magnitudes in the particular case examined; if we make distance the minor premise, size is the conclusion drawn; if we make size the premise, distance is the conclusion. Or, expressed as a proportion, a particular relation,  $s : d$ , is compared with the general ratio,  $S : D$ , by the unconsciously established equation,  $s : d :: S : D$ , with  $s$  as the datum and  $d$  as the conclusion, or *vice versa*. Practically, we carry out this proportion every day, more or less exactly, in our visual judgments. In any given case we form a cerebral minor premise by observation of a particular magnitude; we refer that premise to the standard of past similar experience as its cerebral major premise, and by inference form our cerebral conclusion. If, for instance, we have formed a cerebral datum from a retinal image cast by an object of known magnitude at an unknown distance, or of unknown magnitude at a known distance, we infer distance from size, or size from distance, more or less accurately according as our personal ratio between size and distance has been more or less accurately formed by previous



training. A man who has been accustomed to express in numbers lengths that he has seen, can put a more exact value upon the size of a house or field than a man who has not been so trained.

We are liable to be more or less grossly deceived by appearances. A man accustomed to hilly landscape will, if set down among mountains, over-estimate size and under-estimate distance. Having thus unconsciously prejudged magnitudes, he is surprised to find that men and cattle are not really smaller than usual, and that distances are much greater than he at first imagined. A hazy atmosphere makes hills look like mountains; a clear atmosphere makes mountains look like hills. A Swiss travelling in Scotland commented on the great size of the cattle; a Scotchman remarked that Swiss cattle were very small. An object seen through a telescope appears nearer than it really is, although on other grounds it is known to be at a great distance; conversely, an object looked at through a reversed telescope appears distant, although from other sensory information we know it to be near.

Instances might be multiplied in which the cerebral representation of a given sensation is more or less altered by unusual sensory premises, or even irresistibly distorted by collateral sensations, but we shall only quote one such instance—that known as the Zöllner illusion. The two lines  $ab$ ,  $ab$ , are parallel; but in

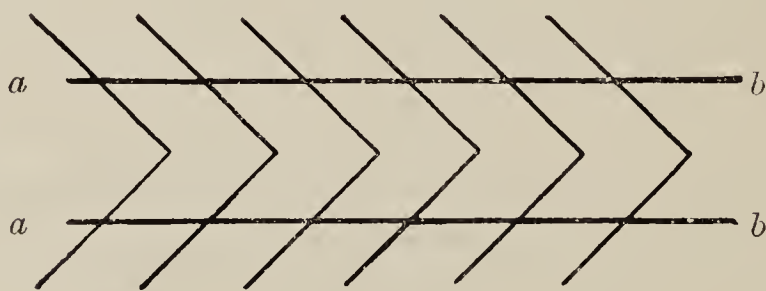


FIG. 301.—ZÖLLNER'S ILLUSION.

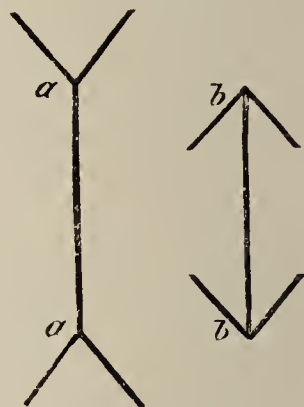


FIG. 302.

consequence of the inclination of the short lines crossing them they appear to diverge from left to right. In fig. 302 the two lines  $aa$ ,  $bb$ , are of equal length, but in consequence of the short diverging or converging limbs,  $aa$  appears longer than  $bb$ .

Finally, we must allude to the fact that the cerebral representation of a sensation or group of sensations may, quite independently of any external distorting cause, be modified by the voluntary attitude and focalisation of attention. The figure

of a beaker may be imagined and 'seen' at will with its open mouth either towards or from the observer; and an actual beaker can equally be 'seen' in either position, especially if one eye be closed, to eliminate collateral items that correct and dissipate the illusion. This elimination is, however, not indispensable; even with both eyes open we may so forcibly focalise attention as to *see* erect a beaker lying on its side, or *vice versa*, and *not see* the lights and shadows and perspective that antagonise the illusion. This simple 'make-believe' experiment is thus illustrative of a very important cerebral custom in the formation of judgments; *i.e.* we see what we attend to, we do not see what we do not attend to. We can wilfully persuade ourselves of a fictitious phenomenon by focalising deceptive data to the exclusion of corrective data; and finally, having formed our fiction, we may consistently use objective data in further distortion of the reality.

This last capacity, which is more or less obviously manifested in every man's daily life, is admirably illustrated by the closer consideration of the beaker experiment. Taking a cylindrical beaker or lamp-glass, placed erect and seen as erect, or placed on its side and seen on its side, the vessel in each case seems cylindrical; but if, by a forcible self-suggestion, the observer 'sees' the erect beaker tilted away from him, the apparent near end looks narrower; or if he sees the overturned beaker as if erect, then the apparent upper diameter seems to be smaller; in each case the really cylindrical object appears to be conical. The observer having formed his false notion, makes further use of the real retinal magnitudes of the image conformably with that notion; in the first fiction the retinal image of the upper end is imagined to be more distant, and therefore appears larger; in the second fiction the further extremity is imagined to be nearer, and therefore appears smaller. The

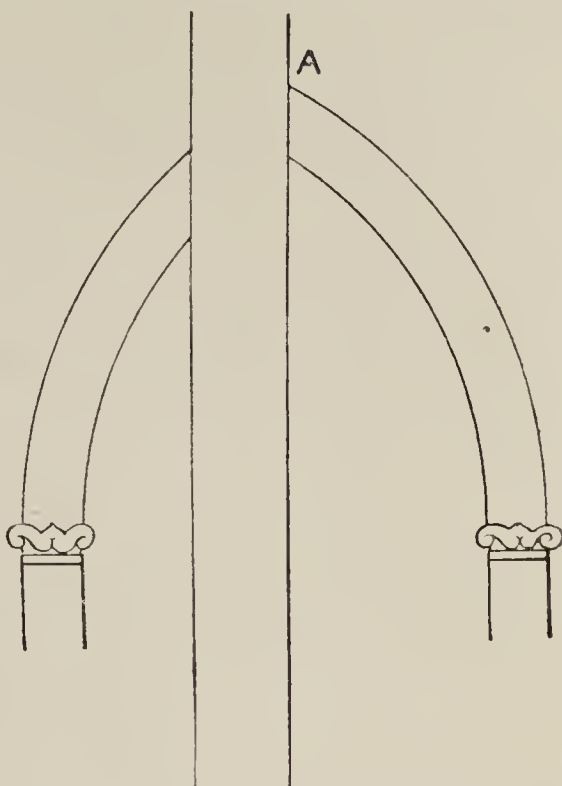


FIG. 303.—APPARENT DISTORTION OF A GOTHIC ARCH VIEWED BEHIND A VERTICAL PILLAR. THE TWO LIMBS OF THE ARCH MEET AT A, BUT APPEAR NOT TO DO SO.

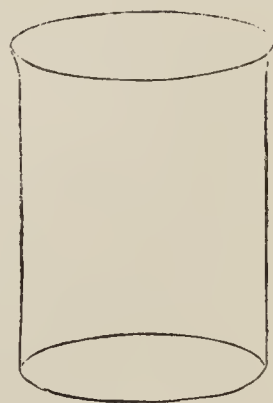


FIG. 304.



great power of this self-deceptive effort may be realised in its fullest extent by holding the beaker in front of the eye and tilting it in various directions; the movements may be 'seen' reversed at will, in spite of the muscular contractions that the observer is making (w.).

There are reasons for thinking that the process of inference, or judgment, as known to us by introspective analysis, and characterised to be an exclusively 'cerebral' function, is also represented in simpler degree in the unconscious operations of subordinate centres, and that we may receive in a supreme centre the *resultant* of impressions already combined and elaborated in a subordinate centre. We have no positive knowledge of the actual seat of formation of particular inferences; they may be superior and inferior cortical, or cortical and sub-cortical; but this ignorance is no bar to the essential conception that such processes may occur at different functional levels. In the case of Zöllner's illusion, the lower-level inference is so strong as to irresistibly elicit the higher-level inference; we *cannot see* the lines parallel, although we may *know* by measurement that they are so. In the series of beaker observations the lower-level inferences are formed from sensations sufficiently ambiguous to allow a higher-level inference to be formed, either rightly or wrongly, and if in that inference *one* of the lower-level components is misjudged, other lower-level components follow suit. With the erect beaker, seen so that the proximal edge appears to be behind the distal edge, that misjudgment of lower in higher inference carries with it misjudgment of the low-level inference from the retinal magnitude of the image. The image of the beaker-mouth is no larger than before, but the beaker is seen tilted away; therefore the low-level inference of size is exaggerated in the high-level compound inference of position *plus* size.

An important means of estimating the distance of objects is afforded by our unconscious appreciation of the difference between the two pictures formed on the two retinae; this factor is especially important in our perception of the third dimension of space, and consequently of the form of solid objects. Length and breadth can be judged of with one eye open as well as with two, but to get the notion of depth we are much assisted by the double sensation of two slightly different pictures. An ordinary landscape, looked at with both eyes open, appears much more extensive as regards the third dimension than when it is looked

at with one eye shut. On the contrary, the painting of a landscape appears much flatter when it is looked at with the two eyes than with one only. These differences may easily be understood: the depth of a landscape is most apparent when two different retinal pictures are formed; the want of depth in a picture is most apparent when two identical retinal pictures are formed; it is especially the foreground effects that influence our judgment in the two cases.

In the more delicate discrimination of distance at closer ranges, *e.g.* when the hand is put out to move a piece at chess, other subjective factors connected with a sense of movement come into play, viz. the degree of convergence of the two visual axes and the amount of accommodation exerted.

The *haploscope* (*ἁπλόος*, single) is a device by which, with parallel visual axes, a single sensation is obtained of two separate visual fields. Whereas in ordinary binocular vision the visual axes converge to a point of fixation, and we have single vision excited by two slightly different retinal pictures, in haploscopic vision the axes are parallel, or nearly so, and we have single vision excited by two altogether different pictures, forming with appropriate objects an illusory sensation. Haploscopic vision

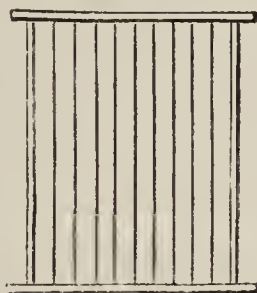


FIG. 305.

To illustrate haploscopic vision. The distance between the bird and cage is here given as rather less than the distance between the two pupils, so that the haploscopic combination can be effected easily, with the visual axes somewhat convergent. The illusion is facilitated by holding a card vertically between the eyes, so as to hide the bird from the left eye and the cage from the right eye.

may be effected by looking through two parallel tubes, and is approximately effected by stereoscopes; it may be simply illustrated by fig. 305, which represents a familiar trick: holding the figure close to the face, and looking beyond it, the bird will appear to slide into the cage as the visual axes become less convergent; we then have in the right field of vision a bird, in the left field of vision a cage, and in consciousness a bird in a cage.

The stereoscope and the telestereoscope are instruments so constructed as to give an exaggerated notion of the third dimension. In the ordinary stereoscope two photographic pictures, taken from slightly different points of view, are looked at, each by one eye, and the visual axes are adjusted until the two pictures become 'identical' on the two retinae, and therefore fused



in consciousness as a single picture, in which 'depth' becomes particularly apparent. This is called a 'stereoscopic effect,' and

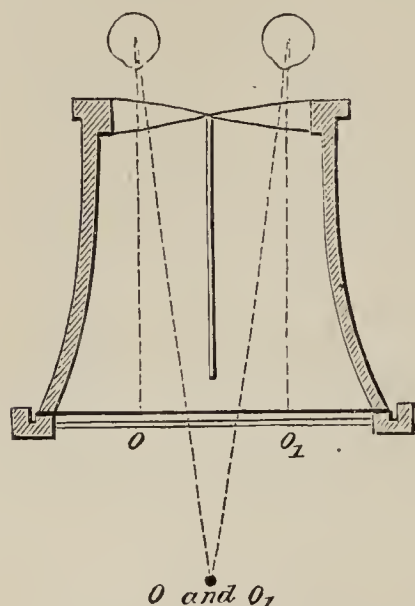


FIG. 306.—STEREOSCOPE.

Two objects,  $o$  and  $o_1$ , are viewed in conjunction.

in the actual instrument the convergence of the visual axes is imitated and exaggerated by two prisms with the thick edge outwards, one in front of each eye. That 'stereoscopic effect' is not due to movements of the eyes is proved by the fact that it is produced with pictures illuminated by the electric spark; no sensible movement of the eye is possible during the illumination thus produced.

If, instead of placing in the stereoscope two photographs prepared *ad hoc*, we examine two identical pictures or prints, their two images give no stereoscopic relief, but combine into a single flat image. If there be the slightest difference between the two prints, this will at once be detected as an irregularity when they are combined by the stereoscope. Two genuine bank-notes can be thus perfectly combined; a genuine and a spurious note cannot.

The effect of the telestereoscope, as will be understood from the accompanying figure, is to exaggerate the difference between the two retinal pictures of a landscape by causing the points of

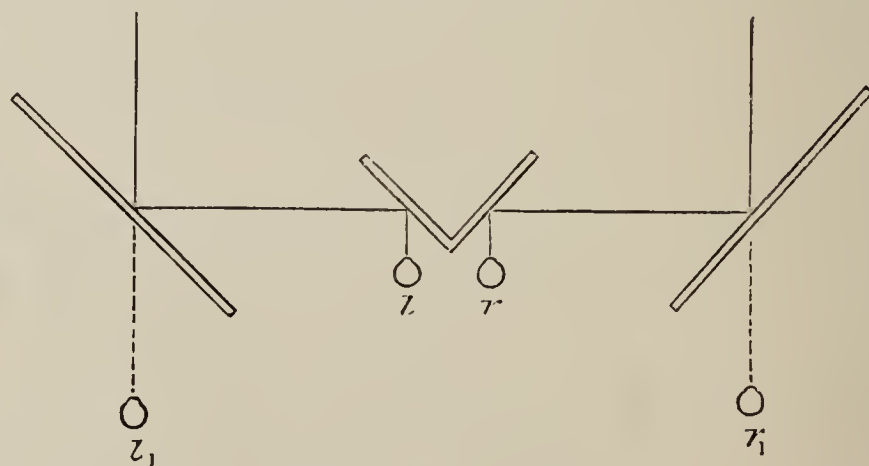


FIG. 307.—TELESTEREOSCOPE.

Two small mirrors in front of the eyes,  $l$   $r$ ; two larger mirrors arranged to reflect images of distant objects upon the smaller mirrors, and thus to the eyes. The effect is as if the eyes were placed at  $l_1$   $r_1$ . (Helmholtz.)

view of the two eyes to be much farther apart than normal. The normal distance,  $l$   $r$ , between the centres of the two eyes being 65 mm., is by the telestereoscope increased to the distance  $l_1$   $r_1$ . If, for instance, this distance be 65 cm., the normal stereo-

scopic effect is proportionately exaggerated, and at the same time objects in the landscape appear reduced in size, producing the effect of stage scenery.

**Contrast.**—Any quality, optical or other, of an object or person causes in the observer an impression that is intensified if the surroundings are of an opposite quality, weakened if they are of a similar quality. Any given quantity is over-estimated when its surroundings are unusually small, under-estimated when its surroundings are unusually large. ‘Rest can only be enjoyed after labour; sound, to be heard clearly, must rise out of silence; light is exhibited by darkness, darkness by light; and so on in all things.’ Stated more generally, the value of any given constant is estimated to be  $+$  or  $-$  according as its surroundings are  $-$  or  $+$ . A person of average height appears taller among short people, shorter among tall people. If one finger dipped in hot water, and another finger dipped in cold water, be both simultaneously transferred to tepid water, the first finger will feel cold, the second warm; these are phenomena of successive contrast, analogous with the negative after-effects of visual excitation.

Formally expressed, if  $A : B$  is the personal ratio or major premise,  $a : b$  the minor premise formed by an object in its surroundings, then the object  $a$  varies inversely as the surroundings  $b$ ; if by the focussing of attention  $b$  is made the premise,  $a$  becomes the conclusion, and *vice versa*.

The theory of contrast has long been a vexed question, and has been debated most closely in the case of simultaneous colour-contrast (p. 460). The question at issue is very similar to that stated on p. 554 with reference to the Weber-Fechner ratio between stimulus and sensation, and as in that case, so in this, a curt, and therefore dogmatic, verdict would be of no value. Weighty names are subscribed to rival theories; Helmholtz, as the chief upholder of the ‘*psychological theory*,’ represents contrast effects as being due to an error of judgment; Hering, as the chief upholder of the ‘*physiological theory*,’ represents contrast effects as being due to the up-and-down chemical modifications of a hypothetical retino-cerebral ‘vision-stuff.’ In the psychological theory, a transforming factor is presumed to occur between sensificatory change and sensation. Helmholtz holds that material sensificatory data do not directly modify each other, but that in the meta-physiological process of inference sensations are modified by other sensations within the field of perception. In the physiological theory, a



direct material modification of sensificatory data by each other is presumed to occur ; Hering urges that each such datum diffuses in the retino-cerebral organ beyond its precise locus of incidence, and directly modifies contiguous sensificatory data, and the observations of Charpentier, alluded to on p. 457, are confirmatory of this mode of view. According to Hering's mode of expression, an opposite or assimilatory change (black, blue, or green) is induced in the retino-cerebral area surrounding a given spot or area in which a dissimilatory change (white, yellow, or red) is provoked—and *vice versa*.

One of the experiments upon which the Helmholtz-Hering debate has especially turned is that already alluded to on p. 461 as Meyer's experiment.

Helmholtz's interpretation of that experiment is as follows : 'If the background is green, the covering-paper looks greenish. If the latter covers the grey scrap without interruption, we fancy that a grey object is seen through a greenish veil ; such an object to look white must be rose-coloured.'

But, says Hering—and the statement is easily verified—a chessboard pattern of grey and green squares, covered with tissue-paper, over which a black paper with a square opening is placed, gives a similar if less pronounced effect ; a person coming fresh to it sees no covering-paper, but merely a pattern composed of greenish and reddish squares. His view is that the contrast effect depends upon the fact that any excitatory disintegration of a retinal area is accompanied by a reverse alteration or integration of the surrounding neighbourhood, or *vice versa*, causing a light halo round a dark patch, red round green, green round red, blue round yellow, yellow round blue.

Hering's fundamental experiment in support of this position, in one of its simplest forms, is as follows : Two V-shaped pieces of grey paper are placed upon white and black backgrounds, as shown in fig. 308. By contrast the V on the dark half looks lighter than the V on the light half. The point of fixation is steadily looked at for half a minute to a minute ; the gaze is then removed to a point of fixation in a uniform surface. The previously dark half of the field looks light as compared with the previously light half ; the after-image of the previously lighter V looks darker than that of the previously darker V ; and even when the contrast between the backgrounds has faded off, the contrast between the two V's still remains perceptible. This after-difference must be due to localised change ; it cannot be

due to false judgments between each V and the corresponding background after-effects. The experiment can be varied with similar effects, to which similar reasoning is applicable, by

taking coloured strips of paper on coloured backgrounds, *e.g.* yellow and blue strips on blue and yellow grounds respectively.

The after-images of the strips contrast, even after the background after-images have ceased to contrast. We may even arrange matters so as to obtain a strong simultaneous contrast between two areas in which contrast-colours are produced by coloured fields that are themselves in much weaker contrast: A white surface in front of the two eyes, with a red glass in front of one eye and a blue glass in front of the other, is red or blue to each eye respectively; a black strip on that surface, seen by both eyes open but not focussed for the strip, forms images on non-corresponding parts of the two retinae, and the two apparent strips are in strong contrast, while the surrounding areas appear respectively scarcely blue or scarcely red, and in much less striking contrast than the strips. (Fig. 309.)

We may make Meyer's experiment, in a form unfavourable to Helmholtz's interpretation, by substituting for the grey paper a piece of the same tissue-paper that is to cover the whole; without the cover the field is green and the patch greenish, with the cover the field appears greenish and the patch rosy, although we *know* that the patch is not white, but greenish. On the other hand, we may give to the experiment a form that is very favourable to the Helmholtz view, thus: A narrow strip of grey paper is placed on a white field; two rectangular pieces of coloured paper are fixed at each side of the strip, bordering

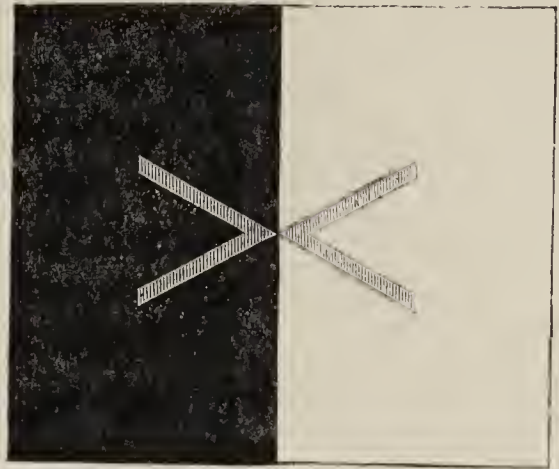


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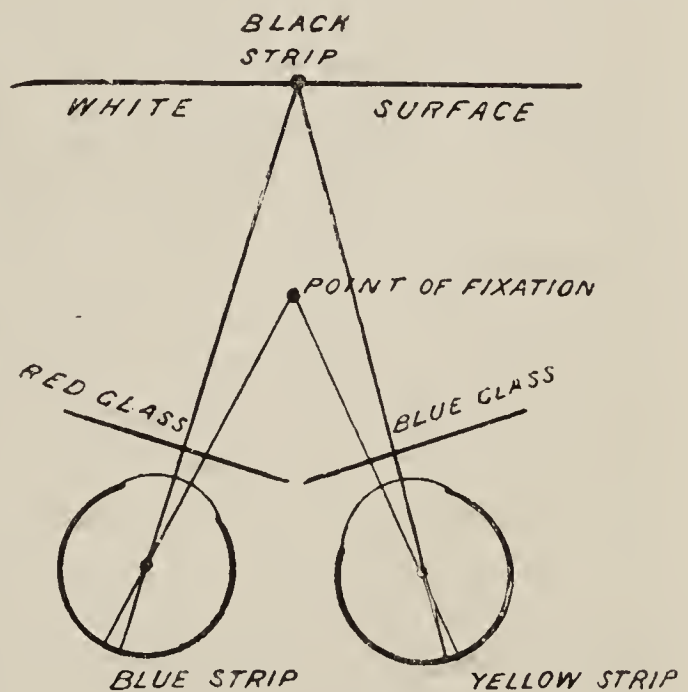


FIG. 309.

strip of grey paper is placed on a white field; two rectangular pieces of coloured paper are fixed at each side of the strip, bordering



upon the middle portion of its length; the whole is covered with tissue-paper. The entire length of the grey strip appears of uniform greyness, or very faintly tinged; but if the coloured rectangle is completed by white or black lines crossing the grey strip, the portion enclosed by these lines assumes the complementary tint, while the portions above and below do not, but, if anything, are slightly tinged in the same sense as the coloured paper (w.).

In sum, what are we to believe? Dogmatic faith is out of the question, and the case is one in which we may fitly anticipate upon a remark below, to the effect that a judgment (or opinion) is the resultant of a synthetic process in which the major premise is the personal ratio of the thinker. It is a truism that psychological phenomena are the subjective symptoms of material sensificatory changes; but it is also an undoubted fact that sensificatory data, while remaining objectively constant, can give rise to sensations that are larger, smaller, positive or negative, according as they are accompanied by other sensations that are relatively smaller, larger, negative or positive.

The term 'judgment' belongs to psychology rather than to physiology, but in any account of the functions of the brain it is not possible to escape the use of terms bordering upon, or actually belonging to, the domain of psychology. We may take this opportunity of recognising one or two of the instances where this trespass has been unavoidably committed. 'Consciousness,' 'judgment,' 'sensation,' 'attention,' 'memory,' are the principal terms thus involved; and although we shall fail to define them strictly, we shall with profit recognise the sense in which each term has been used in the present discussion. *Consciousness* is the subjective experience of each individual; we cannot physically describe the nature of the phenomena that it indicates; we simply paraphrase the term by saying that consciousness is the subjective aspect of objective material changes in the outside world and in the cerebrum of the conscious agent. *Sensation*—i.e. impression of which the subject is conscious—we have used in the sense of 'felt impression' in consciousness, '*perception*' in the sense of 'compared sensation' in relation with associated sensations. *Memory* is the consciousness of former material changes and sensations; objectively it is conceivable as the sum of material cerebral alterations left by previous sensations—the 'organisation of past experience.' *Attention* and *inattention* are attitudes of consciousness, which is concentrated or not concentrated upon given objects. *Judgment*, in its simplest form, or inference, is the resultant of compared sensations, a consciousness of their identity and of their difference, i.e. of their ratio; in its higher and more compound forms it is

the resultant of compared ratios, *i.e.* of a synthetic process in which the major premise is the personal ratio of the thinker.

**Fatigue.**—Fatigue, in the ordinary sense of the word, is a sensation caused by previous exertion or prolonged excitation, and manifesting itself as blunted sensibility or diminished motility. From the physiological standpoint, motor fatigue is recognised in a diminished energy of motion. We have already, on more than one occasion, referred to the signs and symptoms of motor and of sensory fatigue (pp. 333, 382, 459, 558); we have now to disengage the shares of the total burden borne by central and by peripheral organs in the cases of voluntary action and of sensation. We may at once dismiss the latter; we have no means of separating a peripheral from a central factor in the case of sensory fatigue; all that need be said is, that the minimum excitation and minimum perceptible difference between two stimuli must be larger in a fatigued than in a normal state of any sense-organ. As regards voluntary motor fatigue we can, however, by experimental means separately demonstrate, and to some extent measure, central and peripheral loss of energy. We saw in the section on Muscle and Nerve (1) that nerve is practically inexhaustible, (2) that the exhaustion of the motor end-plate precedes and obstructs exhaustion of muscle submitted to indirect excitation. The problem now before us is therefore limited to an appreciation of the distribution of motor fatigue in cerebral and in peripheral organs. The appreciation cannot be made precisely, but we may obtain definite evidence that peripheral and central organs both participate in the change. The mere sense of fatigue contains no proof at all; the material change might be central or peripheral, or both. But that it is in part peripheral is proved (1) by the beneficial effect of massage, (2) by the smaller amount of energy elicited by direct electrical stimulation after exhaustive voluntary action. That it is in part central is proved by the fact that when cerebral action has ceased to be effective on muscle, electrical stimulation of nerve or of muscle is still provocative of contraction. The following experiment may be quoted in evidence of central fatigue:—If a series of induction shocks is applied to a frog's brain and bulb until the gastrocnemius has ceased to respond, a second series of contractions may be elicited by switching the current to a pair of electrodes applied to the sciatic nerve; if, when the muscle has ceased to respond to



this excitation, the current is switched to electrodes applied to the muscle, a third series of contractions is obtained; from which we learn that maximum action of the superior organ does not elicit maximum action of the subordinate organ—in other words, that central fatigue is limitative of peripheral fatigue—and we may formulate, as a probable conclusion, that the incidence of normal voluntary fatigue is in diminishing gravity from centre to periphery—relatively greatest at the former, relatively least at the latter (w.).

Morphological differences in central nerve-cells, after rest and after action, analogous with the differences between loaded and discharged gland, have been recently described by Hodge, who, contrasting, with identical methods, the ganglion-cells of frogs and cats, before and after prolonged stimulation, and the brain-cells of birds and of bees killed at daybreak and at night-fall, finds marked changes, especially in the cell-nuclei—viz. in the morning a plump nucleus staining faintly with osmic acid, as compared with a shrunken dark-staining nucleus in the case of evening sections.

**Sleep.**—After exertion rest is necessary; expenditure of material during the former is made good by restoration during the latter. The brain conforms to this necessity, and does so in a periodic manner. But sleep is not an exclusive attribute of the brain: it extends in less degree to the subordinate nerve-centres, and is attended with a diminished activity of the periphery itself, as evidenced in the slower respiration, slower pulse, diminished excretion of  $\text{CO}_2$ , diminished secretion of urine.

The state of the cerebral circulation during sleep has been the subject of some debate; opposite opinions have been advanced, to the effect that the brain is anæmic, or that it is congested. Although there is no doubt that in coma—a pathological state similar in some respects to physiological sleep—the cerebral vessels are congested, the observations of Durham on the exposed cerebrum of sleeping dogs, and of Jackson on the retinal vessels of sleeping infants, are to the effect that vessels shrink in sleep, and we may therefore feel reasonably assured that the sleeping brain, in common with other resting organs, receives less blood than in its state of activity. Moreover, Mosso's investigations on exposed human brains afford evidence that the organ becomes more vascular during mental activity, and Horsley has seen the cortex of dogs become more vascular during an epileptoid attack.

On the supervention of sleep external impressions cease to be perceived, consciousness fades, the voluntary muscles relax, the eyelids drop, the eyeballs turn up, the pupils contract. The time at and during which different persons sleep is liable to considerable variations, in accordance with temperament and habit; normal sleep lasts from six to nine hours, and it is best that these should be hours of darkness. An excited brain, unusual and irregular stimuli of any sense, are the usual causes of wakefulness; a moderately fatigued condition of 'mind' and body, quietude or monotony of surroundings, are favourable to the arrival of sleep. The depth of sleep is very variable; in deep sleep consciousness is absent, stimuli of even considerable strength may be insufficient to arouse sensation, the muscles are completely relaxed, the so-called tendon-reflexes are in abeyance;

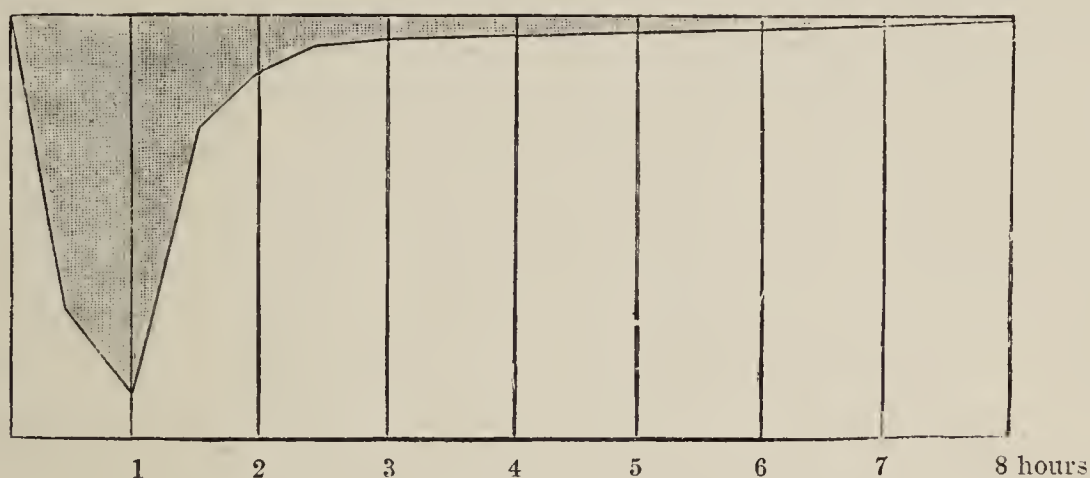


FIG. 310.

To illustrate the depth of sleep, as investigated by the loudness of sound required to wake the subject. (Exner after Kohlschütter.)

the deep sleeper lies dreamless and motionless. All these signs are wanting in sleep that is light, or partial: consciousness is blunted, but not absent; ideas string themselves together in more or less congruous sequence and form dreams, or are aroused by more or less appropriate stimuli, or they actually arouse movements more or less coördinate and purposive, such as are seen in their fullest expression in the actions of sleep-talkers and of sleep-walkers. Normally, a sound sleep is deeper at its commencement than at its end, though abnormally it often happens that a wakeful person sleeps partially and insecurely at first, heavily and deeply later on. The accompanying curve, in which the shaded ordinates represent intensities of sound sufficient to awaken a sleeping person at different periods, gives an idea of the normal depth of sleep from the first to the eighth hour.



**Hypnosis.**—It is not possible at the present day to conclude an account of the action of the central nervous system without reference to the phenomena of hypnosis.

It has long been known that a condition which we now call ‘hypnotic’ is inducible in animals. Kircher’s *experimentum mirabile*, consisting in placing a fowl with its beak to the ground before a chalk-line traced from it, is one of the first on record (1646). Many birds, mammals, amphibia, and reptiles are ‘hypnotised’ by simply turning them on their backs, and a screaming baby can often be thus suddenly surprised and quelled. The phenomena of fascination by snakes are of a similar character. Horses are readily ‘hypnotised,’ and can be shod while in this state of fixed attention.

Fixed attention and obedience to stimuli are, in fact, the essential elements of the hypnotic state in man ; the particular nature of the stimulus by which attention is arrested is of quite secondary importance ; staring at a bright object, listening to a monotonous sound, ecclesiastical stimuli, the scissors of a hair-dresser or the brush of a lady’s maid—in short, anything or anybody—may be the effective cause of a state which, if moderate, is called drowsiness, and if excessive, trance. In the normal state of attention, a person is more alive to a particular class of excitants and less so to all others—he is ‘pre-occupied,’ and, as is well known, his pre-occupation may be slight, or moderate, or considerable. In the hypnotic state the cerebral pre-occupation is excessive—the subject may be impervious and non-sensitive, except to certain focalised objects, or persons, or ideas.

In its commonest forms, the hypnotic state presents many features of resemblance with normal sleep : a hypnotised person appears to be asleep, and in common language is said to be so ; but it is of great importance to know that the most complete hypnosis may occur without any of the superficial signs of sleep (Bernheim). An ordinary person in natural sleep exhibits very slight differences of excitability towards the items of his environment : he is, perhaps, more easily awakened by the sound of his own name than by other sounds. A professional person while asleep is more susceptible to his own professional stimuli than to any others—an exhausted waiter wakes to the whispered call of ‘waiter !’—the slightest wail of her infant wakes its mother. A hypnotised person exhibits inequality of excitability pushed to extremes ; one prescribed kind of stimulation may then act

upon him while all others remain without effect ; he (or she) does or feels whatever the operator of the moment tells him to do or feel, and he cannot do nor feel whatever he is told he cannot do nor feel. Yet it may be easily proved that the stimuli which were unfelt during the hypnotic state did actually produce their ordinary sensificatory effect, and were in reality felt ; *e.g.* a patient on awakening from a state during which he had been made incapable of seeing or incapable of hearing, may describe what had been shown to him during that state, or answer questions to which he had been deaf, but which his very answers prove that he had really heard. It is paradoxical, but nevertheless true, to say that the subject of such false blindness or false deafness, which may be limited to any given portion of his environment, must first recognise the sights and sounds to which he believes himself to be, and therefore is, blind and deaf. A subject told that he cannot see or hear things to his right side must recognise that they are to that side before he is ' blind ' or ' deaf ' to them. Fraud is, of course, possible in this as in other matters ; but, apart from any fraud, a hypnotised person can be at the same time blind and not blind, deaf and not deaf—an apparent contradiction of fact that is only very imperfectly characterised by saying that he is *psychically* deaf and blind, while *physiologically* not deaf and not blind.

An ordinary sleeping person is to a certain extent an automaton, reacting as stimulated ; judgment and discrimination are in abeyance. A completely hypnotised person is, in the fullest sense of the word, an automaton, reacting as stimulated by the operator ; the stimulation or ' *suggestion* ' arouses in the brain of the subject an ' *idea* ' with crude and startling distinctness, unmodified by any scepticism or criticism, and produces with fatal certainty action appropriate to idea. A hypnotised patient believes everything his temporary ' master ' tells him, and carries out every order he receives, moving in obedience to suggested ideas or provoked hallucinations. These have only a subjective existence, as is recognisable by the fact that the apparent object is not modified in appearance by optical instruments : *e.g.* an imaginary blot on a card is not doubled by a prism (unless, indeed, there should be real marks visible and doubled) ; a disc imagined as half red, half blue, is not seen purple when rotated (unless the subject knows that purple is the resultant of red and blue). We recognise in the provoked hallucinations of a hypnotised subject an exaggeration of what normally



occurs in a more or less marked degree in everyday life, not merely in the reflex reactions of the medullary axis, but in those of conscious imagination aroused by real objects. The ideas of things in the mind of the most exact observer are no faithful copy of things in themselves, and in the consciousness of a careless observer depart in most extravagant ways from the actualities as they appear to a majority of normal men; in an insane person, or in a hypnotised person, the idea and the thing diverge to such an extent that the former becomes characterised as a hallucination or a delusion.

A child is 'impulsive,' reacts upon the suggestions of the moment—in a word, is more automaton-like than a reflective self-controlled adult. Religious ecstasy, lovers' imagination, ordinary dream chains, somnambulism, the deceptions produced by the distracting manœuvres of a juggler, offer more or less familiar instances of that concentration or predominance of a train of ideas—to the effacement of other accessory or modifying ideas and sensations—which in extreme degree is characteristic of the hypnotic state. Education is not only instruction, it is suggestion working upon brains more or less predisposed to reception, more or less pre-occupied by the effects of previous suggestions; and the influence of some persons upon the beliefs and conduct of others is an everyday instance of physiological hypnotism, the actual result depending upon two factors—upon the impressiveness of the operators, upon the susceptibility of the subjects. A hypnotised person is in a state of '*suggestibility*,' or unsceptical credulity, which is a retrogression towards a primitive state, and an exaggeration of that working credulity of everyday life which enables us to acquiesce in and act upon simple statements without constantly exacting argument, or evidence, or proof. We naturally—in the absence of stronger reason to the contrary—believe what we are told, and imitate the actions of other people. Suggestions to ideas and to actions have more or less pronounced effects on persons of different temperament; they have an exaggerated or forced effect upon hypnotised subjects during the passive state, or even, it may be, in their ordinary awakened condition. All men are more or less automata; hypnotised subjects are excessively or completely automata.

The influence of mind on body—using this expression in the ordinary undefined, yet sufficiently indicative sense—offers instances which, although not embraced by the term '*hypnosis*,'

bear, however, some relation to its phenomena, there being this point of similarity between them—that extra-ordinary body effects are produced in both cases by concentrated mental attitudes. A person having swallowed a bread pill in the belief that he has taken a purgative, will very possibly be purged ; a person having received a hypodermic injection of water in the belief that morphine has been used, will very possibly go to sleep ; railway-litigation-shock is mainly a disease by imagination, but none the less a very real disease. Far more surprising and, to all appearance, well-authenticated instances might be quoted (*e.g.* stigmata, faith-cures, &c.), but with marvels *qua* marvels we are not concerned ; the simple instances just given are quite sufficient to bring home to us the fact that ‘state of mind’ is a potent factor in state of body, and that the first aim of the physician should be to establish in his patient the idea of recovery.

Two great practical lessons are the immediate outcome of any intelligent study of hypnotic phenomena ; the first is the lesson of self-distrust, the second is the lesson of self-confidence.

Every man’s brain is more accessible to certain kinds of ideas than to certain other kinds ; all men, even the most judicial, form opinions and convictions upon data selected by self-suggestion, coloured and refracted by their own mental media. We see and hear what we look or listen for, we believe what we wish for, and each man, outside the province of his own tastes and sympathies, is in varying degree blind and deaf to actual phenomena ; what he dislikes he disbelieves. A full and real apprehension that this is a necessary attribute of cerebral action cannot but be a corrective of that ‘cocksureness’ which is one of the blind-alleys in which men’s minds are apt to become blocked, and give practical value to the sentiment more often expressed than felt—‘*You may be right ; I may be wrong.*’

It is no less important—and for the physician it is immediately and specifically important—to fully realise the power of suggestion on men’s minds and bodies. Like breeds like ; self-distrust in the guide gives rise to want of confidence in the guided ; self-confidence in the leader raises confidence in the follower. Nothing ensures success more than the confident expectation of success ; the fear of failure often brings failure as its direct consequence. The physician who knows and trusts his own resources will carry his patient through illness, where another man will vacillate and despond, and by his face or manner suggest failure and hasten death.





## APPENDIX

### I. ORIGIN AND NUTRITION OF THE EMBRYO

THE OVUM.	Maturation . polar globules, female pro-nucleus	} germinal vesicle.
	Fertilisation . spermatozoon, male pro-nucleus	
	Segmentation.	
	Organisation . The three primary layers, their fate and functions in the adult.	
<i>Hypoblast</i> .	The digestive tract.	} The allantois and placenta.
<i>Mesoblast</i> .	The vascular system. First, or vitelline.	
„	Second, or allantoic. Vascular changes at birth.	
<i>Epiblast</i> .	The nervous system.	} The amnion.

**Experimental physiology of the embryo.**—Composition of a fresh hen's egg and of a newly hatched chick. Respiration within the egg. Origin of hæmoglobin.

Respiration of the mammalian embryo. Blood-gas analyses. Blood-pressure and blood-flow in the umbilical vessels. Hæmoglobin. Heat-production. The renal function; indigo-carmin experiments. The digestive ferments. The hepatic function; meconium. Glycogenesis. Structure and properties of the chick's heart *in ovo*. Reflex movements of the embryo.

WE do not undertake to give even a summary description of the anatomical development of particular parts, organs, and tissues. However essential such descriptions may be to the comprehension of adult malformations of medical as well as of surgical interest, we regard them as being outside physiological bounds, and to be learned far more thoroughly and easily in their appropriate place. Nor shall we enter upon any description of the mechanism of coitus or of the phenomena of menstruation and parturition; we may, indeed, point to the culminating event of coitus—viz. 'ejaculation'—as affording a well-marked example of spinal reflex discharge in consequence of summated stimuli (p. 497), and we may mention the fact that parturition itself can be effected as a reflex act from the lumbar portion of the cord separated from the brain by transverse division; but the detailed description of parturition and of menstruation belongs properly to the province of



obstetrics ; the only points which may, perhaps, be mentioned here, as being of physiological interest, are that each menstrual flux is probably accompanied by the discharge of an ovum, that the most frequent seat of fecundation of ovum by spermatozoon is probably the upper part of a Fallopian tube, and that, according to the somewhat limited range of trustworthy statistics available, impregnation appears to have occurred most frequently during the first week after a menstrual period.

There are, however, several organs, the functions of which are accessible to study in embryonic life, and general ideas relating to nutritive functions illustrated in the development of the embryo, as well as various anatomical questions immediately bearing upon physiological doctrine, that demand attention from a physiological standpoint. These we shall consider, and in so doing shall find it necessary to recognise where we are, and how certain embryonic conditions have been reached.

The entire human body is originally derived from a single female cell, the *ovum* or germ-cell, modified by the action of a male cell, the *spermatozoon*. The process of fertilisation consists in the conjugation of these two cells or of portions of them, and is preceded by certain preparatory changes occurring in the ovum prior to the advent of the spermatozoon, collectively termed *maturation*, viz. an extrusion of part of its nucleus in the form of the first and second 'polar globules,' the remainder of the nucleus being termed the 'female pro-nucleus.' Maturation of the ovum having been accomplished, *fertilisation* proper commences by penetration of a spermatozoon ; the head of the spermatozoon enlarges, forms the 'male pro-nucleus,' and fuses with the female pro-nucleus, forming a new nucleus, the true germinal vesicle ; the nucleated cell is now fertilised, and capable of undergoing cell-multiplication by cell-division and cell-growth. The next stage is *segmentation*. The single cell, with its nucleus formed from male and female pro-nuclei, divides into two cells, each of these into two, and so on, giving rise to a growing mass of multiplying cells, 2, 4, 8, 16, 32, 64, &c. In mammals, the entire mass of the cell is involved in this process ; in birds, it is confined to a small portion of the much larger mass that forms the yolk of an egg.

We may at this point attend to the differences of structure, development, and nutrition between the two classes of ova, the fate of which most concerns or instructs us, *i.e.* that of a bird, and that of a mammal. A hen's egg comprises the ovum proper, or yolk, and accessory envelopes of albumen and of mineral matter ; if it be a fertilised egg, it will after twenty-one days' incubation contain a bird, which has replaced the originally white and yellow materials. These materials were in chief part the food-store, at the expense of which an originally minute portion of protoplasm has grown, multiplied and developed. A human ovum is a small cell ( $\frac{1}{5}$  mm. in diameter), comprising little more than its own protoplasm ; its yolk is in very scanty amount, it has no sur-

rounding albuminous provision. The whole cell divides and subdivides if fertilised, and not having in itself any considerable store of food, its nourishment and augmentation depend upon the mother-organism, at first by simple imbibition, later by nutritive and excretory osmosis through a special organ—the placenta—which is its respiratory and alimentary and excretory organ.

The first step in the development of the mammalian ovum is cell-multiplication; the first step in organisation is the regular arrangement of its now numerous cells as a vesicle, the blastodermic vesicle, with an accumulation of small granular cells at one pole—the future blastoderm. The first step in differentiation is the distinction visible between upper or outer and lower or inner ranks of cells at this spot, soon followed by an obvious triple division into upper or *epiblastic* lower or *hypoblastic*, and middle or *mesoblastic* layers. This tripartite distinction is fundamental; the different tissues and organs in adult life are traceable back to these three stocks respectively, and already in the three ancestral membranes we may recognise in the germ a differentiation of function that is preserved throughout life. The upper or external layer is neural; it will form epithelium—the general epithelium of the skin, the special epithelium of sense-organs, and, by involution, the central axis of the cerebrum and spinal cord. The lower or internal layer, lying on the yolk, is nutritive; it will form the digestive and absorbent epithelium of the intestine and intestinal glands. The middle layer is the massive working layer, the future bone and muscle of the body, and at a very early stage it shows tokens of the depuratory and excretory functions which are among its functions in after-life; it is the nidus of origin of the entire vascular system and of the blood, and one of the earliest organs to appear in it is the Wolffian body, which forms in the adult the essential parts of the renal and generative organs—of the kidney, testicle, and ovary.

The dorsal or epiblastic surface of the blastoderm rises in two longitudinal ridges which, coming together and coalescing in the middle line, form the medullary tube or axis. The anterior, posterior, and lateral margins of the blastoderm turn downwards and converge, enclosing a cavity, the visceral tube, between which and the yolk-vesicle a communication long remains as the vitelline duct. In the bird's egg, towards the middle of incubation the vesicle contains the true yolk, surrounded by blastoderm; in the mammal, the 'yolk-vesicle' contains much more than the original yolk; it is of new formation, consisting of all that part of the blastodermic vesicle which is not directly involved by the forming embryo, and is termed the umbilical vesicle.

The ventral or hypoblastic surface, in consequence of the folding downwards of the entire blastoderm, soon forms a widely-open ditch, terminating in two cæca—foregut and hindgut—which at a later stage, when communication is established with the exterior by the intrusion



of the stomodæum and proctodæum, become œsophagus and rectum. The intermediate portion, becoming tubular, lengthens and curls up, dilates in some parts, and pushes out processes at others, forming finally the rest of the intestinal canal and its accessory digestive glands (liver, pancreas). A protrusion from the hindgut forms the allantois, the dilated stalk of which persists in after-life as the urinary bladder. The anterior portion or foregut likewise buds forth processes, some of which form temporary communication with the exterior, as the pharyngeal clefts, while others ultimately give rise to permanent glands (thymus, thyroid), which become ductless by obliteration of their original stalks, and others finally form the salivary glands and the most important of adult nutritive and excretory organs—the lungs. The only so-called ‘glands’ that do not originally spring from the hypoblast are the supra-renals, which are developed in association with the sympathetic system from mesoblastic, and probably also from epiblastic, tissue; and the spleen, which is an appendage of the vascular system. The cutaneous glands form also an exception to the rule; they are of epiblastic origin.

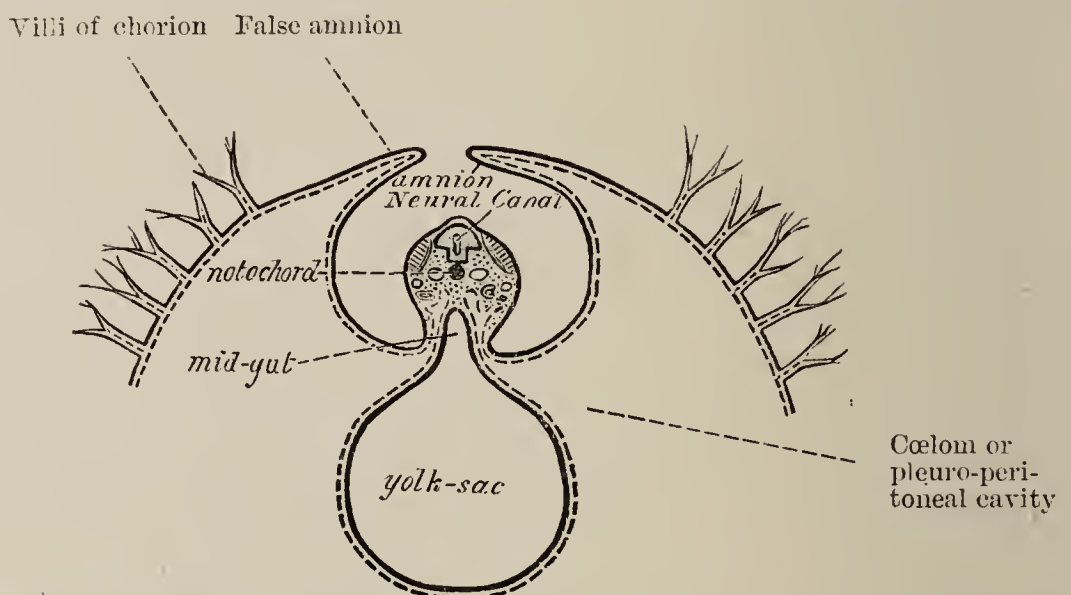


FIG. 311.—DIAGRAMMATIC TRANSVERSE SECTION OF EMBRYO AND OF ITS ANNEXES.

The mesoblast is indicated by the broken line. (Quain's Anatomy.)

With regard to the middle or mesoblastic layer, the most important points to be understood, in addition to those mentioned above, are, its relations to the fœtal membranes, and to the two layers above and below it. Its lateral plates, as distinguished from the axial mass, are themselves distinguishable as somatic and splanchnic; the former or upper layer, conjoined with the epiblast, forms the body-wall and the dorsal membrane or amnion, which surrounds and encloses the embryo, as will presently be described; the latter or lower layer, conjoined with the hypoblast, forms the visceral wall and the secondary ventral vesicle or allantois, which will form the temporary organ of fœtal nutrition—the placenta.

Early in the existence of the embryo, folds arise around it, formed from the dorsal or somatic layer, which finally coalesce dorsally and

form an enclosing membrane, the *amnion*. Of the double layer by which the enclosure is effected, the outer or false amnion fuses with the general covering or chorion; the inner or true amnion alone remains as a distinct membrane; ultimately the embryo floats free in its cavity, which is filled with fluid; its only attachment to the mother is by the umbilical cord, which, as we are about to describe, is formed by the allantois. The main use of the amniotic fluid is mechanical, affording protection to mother and child from sudden movements of either; it is a dilute fluid, with a little albumin and a trace of urea; at term it amounts to between 500 and 1,000 c.c. It can scarcely be regarded as a nourishment-fluid, or as an excretion-fluid; the embryo gets proteid and gets rid of urea by way of the placenta.

A more important structure, physiologically, is the *allantois*; budding forth from the hindgut, it insinuates itself between the two layers of the amnion, and coalesces with the outer layer. It is pre-eminently

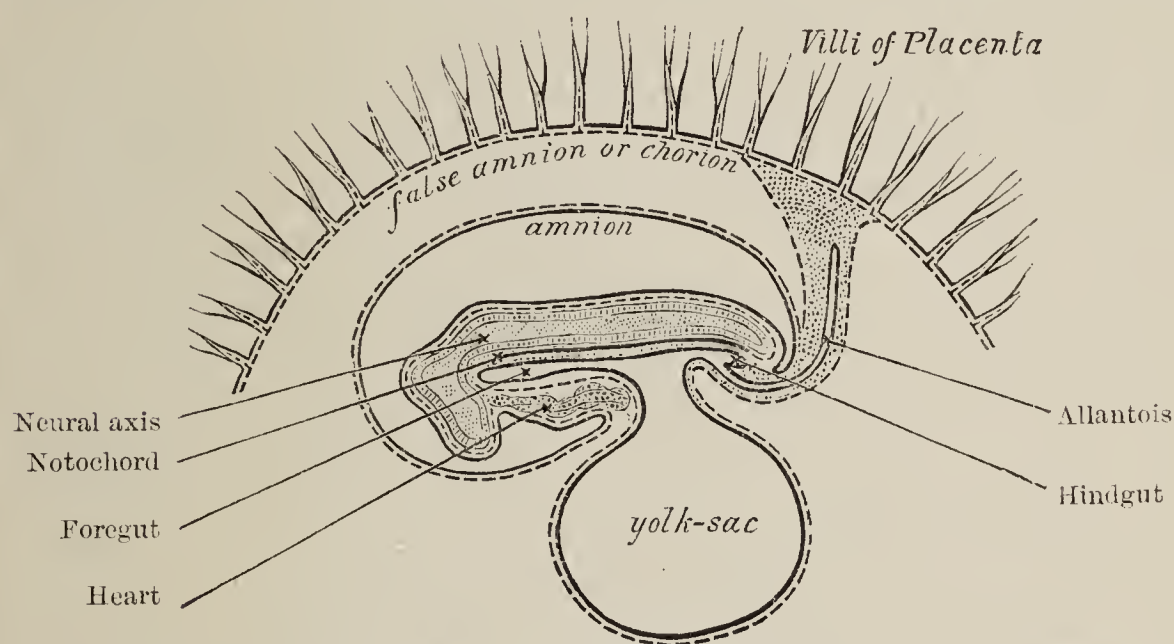


FIG. 312.—DIAGRAMMATIC LONGITUDINAL SECTION OF EMBRYO AND OF ITS ANNEXES.

The mesoblast is indicated by the broken line. (Quain's Anatomy.)

a vascular organ; the ramifications of the allantoic vessels occupy the embryonic villi, which, after an early stage as a uniform shaggy covering, ultimately dwindle at parts and enormously increase at one spot in correlation with increase of the uterine lining. The *placenta* is thus established as the temporary organ through the medium of which everything needed by the growing embryo is obtained from the mother, and everything produced by its metabolism discharged. The organ must therefore transmit proteid, fat, carbohydrate, water, salts, and oxygen from maternal to foetal blood, carbon dioxide and urea from foetal to maternal blood. The weight of the placenta ranges from 500 to 1,000 grms., and, as a general rule, the larger the child, the larger the organ.

*The embryonic vascular systems.*—The first circulation of blood established in connection with the embryo is that of the yolk-sac by the *vitelline* arteries and veins; the ramifications form the *area vascu-*



*losa*, resting on the surface of the yolk; it is the vehicle by which food is conveyed from yolk to embryo. The vitelline circulation is a temporary organ, forming, increasing, and then dwindling away with the gradual consumption of the yolk, and subsequently replaced by the allantoic circulation; in the bird's egg it is comparatively far more prominent than in the mammalian ovum, in which the chief nutrition is from the mother by way of the placenta.

The second circulation of blood is established by the allantois, which arises from the hindgut as just described, and increases as the yolk-sac decreases; its pedicle of origin becomes the umbilical cord, its vessels the umbilical arteries and veins. Vessels within the embryo, connected with these two extra-embryonic systems, form and alter in correspondence with their alterations. The heart appears, first as a double tube, then as a single sac, which pulsates rhythmically before nerves, or even muscle, or blood-corpuscles have made their appearance. The vitelline blood comes to the embryo from the '*area vasculosa*' of the yolk-sac by the vitelline veins, and returns by the vitelline arteries. Later, when the vitelline has made way for the allantoic system, blood comes from the placenta by the umbilical vein in the arterialised state to the embryo, and returns to the placenta in the venous state by the umbilical arteries. Now—that is when the placental function is fully established—the embryonic vascular system exhibits features of structure in obvious correspondence with physiological conditions. The umbilical vein, bringing oxygenated and otherwise renovated blood, enters the liver, some blood passing straight on by the ductus venosus to the inferior cava, the rest after a portal ramification in the liver. The mixed blood in the cava is sent through the right into the left auricle, being guided through the foramen ovale by the Eustachian valve. From the left auricle it passes to left ventricle and to aorta, thus in main part to the cephalic end of the body, whence it returns by the superior cava to the right auricle, through which it is guided by the Eustachian valve into the right ventricle. From the right ventricle it is sent through the pulmonary trunk, ductus arteriosus, and thoracic aorta, in main part to the placenta by the umbilical arteries, in small part to the caudal end of the body.

Thus, in the foetal circulation we have to distinguish at least four qualities of blood—(1) the purest blood, returning from the placenta, and for the most part passing through the liver; (2) the least pure blood, returning from the inferior extremities; (3) the mixture of these two kinds, passing to the left ventricle and cephalic end; (4) the returning cephalic blood, passing to the caudal end and to the placenta.

The noteworthy features from a physiological standpoint, briefly enumerated, are: (1) the smaller difference between typical venous and typical arterial blood in the foetus than in the adult; (2) the great size and importance of the liver, which exercises its action upon almost

the entire mass of the incoming blood—an action which has its most obvious tokens in the comparatively large amount of oxygen lost by that blood in its transit through the liver, and in the large accumulation of biliary excreta in the meconium formed during foetal life ; (3) the fact that, whereas the head end of the foetus receives mainly new

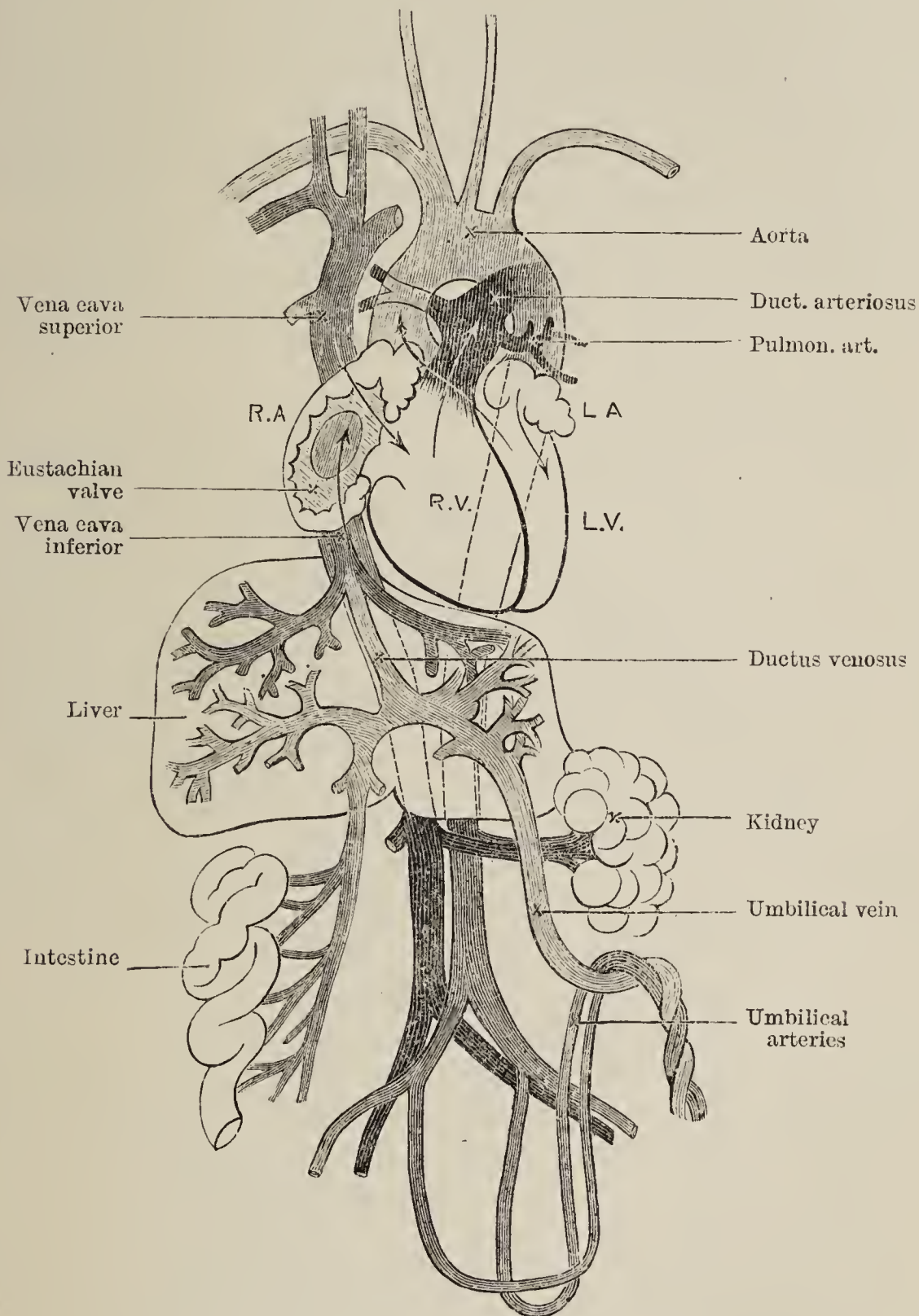


FIG. 313.—THE FŒTAL CIRCULATION. (Quain's Anatomy.)

blood, the tail end receives only second-hand blood ; it will therefore not be matter for surprise that at term the head is far more bulky than the inferior extremities, in comparison with their relative bulk in after-life ; (4) the equal thickness of the right and left ventricles of the foetal heart, in contrast with their inequality in the adult organ ; the equality



and inequality being in obvious correlation with the equality and inequality of resistance to be overcome by the two ventricles in the two states respectively. During the first few days of independent life, in consequence of its separation from the placenta and of the expansion of the pulmonary circulation, the circulatory currents are completely altered; the umbilical blood-flow ceases, and the vessels close, the ductus venosus ceases to be pervious, the pulmonary vessels expand, the ductus arteriosus shrinks and becomes impervious; finally, the foramen ovale between the two auricles becomes more and more completely sealed by the adhesion of its valvular membrane to the margin of the orifice. The cardiac chambers and blood-flow are now in their well-known adult form.

*Origin of the nervous system.*—A knowledge of the developmental history of the nervous system is of some assistance to the due comprehension of its functional significance, and we shall therefore allude to one or two leading facts, and give a tabular summary to show the genealogy of the main masses in the adult brain. Originally, the whole cerebro-spinal and sympathetic system is of epidermic or epiblastic source. The outer or upper layer of the embryo grows in, and becomes embayed and cut off by the medullary folds to form an axial dorsal tube. The anterior part of the tube becomes dilated; the dilatation, being incompletely chambered by transverse constrictions, forms the three primary cerebral vesicles, fore-brain, mid-brain, and hind-brain. The fore-brain pushes out a lateral pair of hollow processes, which form the optic vesicles and become the retina and optic nerves, and an anterior pair of vesicles, which become the hemispheres of the cerebrum. The nerves, cranial, spinal, and sympathetic, arise as outgrowths budding forth from the medullary grey axis and extending with the progressive extension of the periphery. The correspondence between parts of the embryonic brain and parts of the adult brain is as follows:

I. *Fore-brain*=3rd ventricle and optic thalamus.

its lateral vesicles=optic nerves and retinae.

its anterior vesicles=hemispheres and corpora striata; olfactory lobes.

II. *Mid-brain*=iter a tertio ad quartum ventriculum, corpora quadrigemina and crura cerebri.

III. *Hind-brain*=4th ventricle, cerebellum, pons, spinal bulb.

The internal surface, lined by ciliated epithelium, is recognisable in the adult as the central canal of the cord and ventricles of the brain. The surrounding grey matter forms the central or medullary grey matter of the bulb and cord.

The most striking features, from a physiological standpoint, are (1) that the medullary grey axis forming the grey matter of the spinal bulb and cord, is the original stem from which nerves take their source and the higher centres expand; (2) the excessive development of the cerebral hemispheres, which in man grow above and overshadow,

physiologically as well as anatomically, the bulbar end of the medullary axis, and cause it to occupy a comparatively unobtrusive position, under cover as well as under control.

The special sense-organs are of epiblastic source. Tactile organs are, and remain, superficial. Auditory and olfactory organs arise by recession and burrowing of the superficial epiblast; the nose remains freely open, the internal ear is cut off and enclosed by secondary mesoblastic development. The visual organ, on the other hand, although in its essential parts epiblastic, is completed in structure by mesoblastic contributions; the most essential part of the organ—the retina—arises as a hollow process from the fore-brain, itself of epiblastic origin; while the lens is produced by inward recession of the epiblast, and received by the optic vesicle, which becomes folded round the ingrowing mass as a double layer, leaving, however, a gap at its inferior margin.

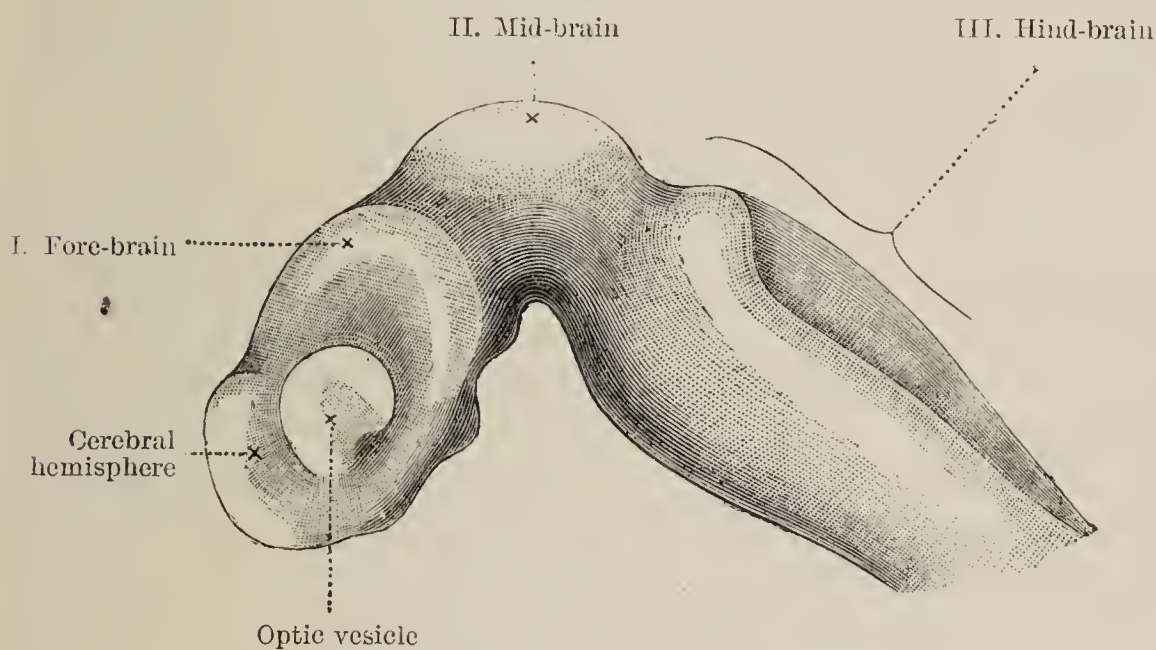


FIG. 314.—THE FETAL BRAIN. (Quain's Anatomy after His.)

This gap, but for which the lens and vesicle might be correctly represented as head and double-layered nightcap, is the choroidal fissure, through which mesoblastic tissue subsequently reaches the interior of the eyeball, and forms its internal vascular and connective-tissue constituents—vitreous body, arteria centralis, iris, and ciliary muscle. The external vascular and fibrous coats—sclerotic, choroid, cornea—are formed from the surrounding mesoblast.

The development of nerve-tracts has already been considered in some detail in connection with the spinal cord and with the regeneration of divided nerves; to the sequence—pale fibre, fine medullated fibre, coarse medullated fibre—we may prefix the still earlier stage of *neuroblast*, which is described by His as the earliest stage of nerve-cell and dependent nerve-fibre, as distinguished from the *spongioblast*, which is the origin of the neuroglia cell and network, forming the supporting tissue of grey matter.



*Experimental Physiology of the Embryo*

From the physiological anatomy of the embryo many general principles regarding its nutrition may legitimately be inferred. We now propose to briefly review the chief points actually verified by observation and experiment, confirming and amplifying such inferences.

*Chemical composition of hen's egg at beginning and end of incubation.*—A fresh egg weighs about 50 grms. and has the following composition :

Shell	5								
White	30	{	Proteids (albumin and globulin)	.	3	}			
			Fats, &c.	.	0.3				
			Salts	.	0.1				
			Water	.	26.6				
Yolk	15	{	Proteids	.	1.7	}	Solids	9.4	
			Fats	.	3		Water	35.6	
			Lecithin	.	1		Total	45 grms.	
			Vitellin, nuclein, glycogen	.	0.2				
			Salts	.	0.1				
			Water	.	9				

The provision in the fresh egg is nearly 5 grms. proteid and over 3 grms. fat; of carbohydrate, only traces, it being formed (? from proteid) in the course of incubation. During the twenty-one days' incubation the shell undergoes no change, and there is no difference appreciable between the salts of a fresh egg (without shell) and of a newly hatched embryo; *i.e.* the embryo gets salts from the contents of the egg, and not from its shell. The contents of the egg weighing 45 grms., the embryo chick formed therefrom will weigh about 10 grms. less; the loss consists of water, of carbon dioxide (.5 gm.), and, according to Liebermann, of an undetermined quantity of nitrogen. The newly hatched chick, weighing 35 grms., consists of about 7 grms. of solids and 28 grms. of water.

As regards water, the early embryo contains it in larger proportion than the contents of the egg or than the formed chick, *e.g.* 90 per cent. in the former, as compared with 80 per cent. in the latter two; although during the period of incubation water has been lost from the egg. Fat (in the yolk) forms fatty acid during incubation (Liebermann). The important point to recognise is, that death of tissue is associated with the very beginning of the life of tissue—katabolic effects are manifested at the very outset in association with anabolic phenomena. But not only does a fertilised egg exhibit respiratory phenomena by yielding CO<sub>2</sub>—an unfertilised egg likewise does so; this is token that such an egg is also composed of living matter, although that matter is incapable of spinning the complete cycle. On reference to the table at p. 132, it may be noticed that the respiratory activity of the chick-embryo has about the same value as that of a hibernating animal: if

the shell, which is the porous septum through which the exchange of gases is effected, be varnished, the inhabitant will perish by asphyxia.

*Respiratory activity of mammalian foetus. Blood-gas analysis.*—

The most obvious of the functions of the placenta is respiratory: blood passing from mother to child by the umbilical vein is of brighter colour than blood passing from child to mother by the umbilical arteries. And further insight has been obtained into the process by the gas-analysis of the blood of sheep-embryos, which at full term weigh about as much as human embryos. The following values were obtained by Zuntz and Cohnstein:

Lamb at full term weighing 3.6 kg.	umbilical artery	$\text{CO}_2$	$\text{O}_2$	
		47	2.3 vols.	per 100
"	"	"	"	"
"	umbilical vein	40.5	6.3	"

the rate of blood-flow was about 44 c.c. per minute, *i.e.* carrying 1.76 c.c.  $\text{O}_2$  and 2.86 c.c.  $\text{CO}_2$ , *i.e.* the respiratory activity of the foetus was less than  $\frac{1}{10}$  that of a normal adult sheep.

The *blood-pressure* in the same instance was between 6 and 8 cm. Hg in the artery, and half that value in the vein, thus illustrating the fact that the difference between arterial and venous blood-pressure is much greater in the adult than in the foetus.

An important peculiarity exhibited by foetal blood, and referable to the fact that such blood contains an abundance of active cellular elements (nucleated red corpuscles), is the much greater rapidity of respiratory changes occurring in the blood itself; in a given volume of foetal blood a larger quantity of  $\text{O}_2$  is consumed and of  $\text{CO}_2$  is produced than in an equal quantity of adult blood. Also, in relation to foetal respiration, it should be mentioned that the amount of hæmoglobin in the body is at first much below that of the adult, and that it gradually approximates to it in the course of pregnancy; at half-time the relation of hæmoglobin to body-weight is only  $\frac{1}{700}$ , at full time  $\frac{1}{200}$ , in the adult  $\frac{1}{140}$  (egg and hen); in an early sheep-foetus the hæmoglobin value may be only  $\frac{1}{10}$ th that of the adult; at term it amounts to  $\frac{9}{10}$ . From which we learn that the respiratory function develops slowly. At birth it undergoes a sudden change—in the sheep it is increased tenfold; from which we learn further, that the conditions of active respiration are already present at a time when, for want of excitant, the process itself is still smouldering at low intensity.

The tissues of an embryo produce far less heat than those of the adult; there is no such thing as thermotaxis during embryo-life, the temperature of a foetus is only one or two tenths of a degree above that of the mother; the heat of the embryo is carried off by the uterine circulation, precisely as heat produced by the liver is carried off by the hepatic circulation; this low production of heat is in correspondence with the low respiratory activity and with the tenacity of life exhibited by an embryo deprived of oxygen; an embryo is slowly asphyxiated in utero if the placental circulation is arrested, it is rapidly asphyxiated by



asphyxia of the mother with intact placental circulation; in the former case embryonic tissue slowly consumes the oxygen it possesses, in the latter case the maternal tissue rapidly consumes oxygen and withdraws it from the foetus.

It is probable that the nitrogen metabolism is also of much lower intensity in the foetus than in the adult; to the question as to its mode of elimination by a foetus in utero, it may be answered that although towards the end of the pregnancy the kidneys are evidently beginning to act, it is probable that the placenta is the main organ of urinary excretion during the period of pregnancy. The bladder of a new-born foetus is sometimes empty, sometimes full; the amniotic fluid contains only a small proportion of urea; cases are on record of a foetus born at full term and possessing no kidneys. These are the main facts upon which the answer is based.

The entire nutrition of the foetus after an early period is, in fact, dependent upon the placenta; the amniotic fluid, to which nutritive action has been attributed, may, indeed, slightly contribute in this direction, but in very minor degree; it contains very little albumin and no carbohydrate or fat; whereas the placenta is a highly glycogenic organ, and has also been asserted to contain peptone.

That the amniotic fluid is of maternal rather than of foetal origin is clearly shown by Zuntz's experiments; after injection of indigo-carmin into the vessels of a pregnant sheep, he found the placenta and amniotic fluid coloured blue, without extension of colour to the foetus itself, or even after destruction of the foetus. On the other hand, after direct injection of carmine into the foetus he found its kidneys blued, which was proof that the organs, even if not ordinarily in action, yet are evidently capable of excretory activity.

With regard to the existence and digestive activity of embryonic ferments—ptyalin, pepsin, pancreatin—artificial digestion experiments have furnished no very precise results, either positive or negative; or, rather, both positive and negative results have been reported by different observers. There can be no doubt that the digestive action of foetal glands is, at any rate, much less constant and pronounced than that of adult glands, but it is probable that the ferments do already exist in the foetal organs in their zymogen state.

By far the most important of foetal organs is the liver; beginning at about mid-pregnancy, or even sooner, the excretion of bile and its accumulation in the meconium are unmistakable tokens that important chemical events are taking place in the liver; so also is the fact, alluded to above, that the liver is the chief seat of the oxygen consumption by the foetus.

The foetal pulse can be felt in the umbilical cord for a short time after delivery, but with the closing up of the umbilical artery it vanishes. The first respiration of a new-born infant occurs in response to the combined effect upon the spinal bulb of the increasing vascosity

of the blood following the arrest of the placental respiration, and of cutaneous stimuli by cold air, or it may be by the flip of a wet towel.

In normal parturition, the birth of the child precedes that of the placenta, and after a shorter or longer interval the cord is ligatured and divided. It makes a great difference to the child whether this be done immediately or some time after birth; in the latter case uterine pressure squeezes blood from placenta to child, and on the average the difference so effected is from 50 to 100 c.c. A normal child of 3 kg. should have 300 c.c. of blood, a child after immediate ligature may have as little as 200 c.c. The fact has been verified on the human subject; a child placed on a balance shortly after delivery, and left connected with the placenta in utero, gradually increases in weight by about 50 grammes.

The *chick's heart*, from the fourth day of incubation onwards, is capable of being submitted to certain observations and experiments; the contraction, which is at first at very irregular intervals, sweeps over the tube in a peristaltic manner from the venous to the arterial end, and by means of photography its rate of progress has been determined to be 5 to 10 mm. per sec. (Fano). Placed in a watch-glass in a few drops of fluid, it forms an exceedingly sensitive reagent to the action of nutrient or non-nutrient fluids; the beats die out in salt solution, and may be renewed by serum-albumin; examined in a gas-chamber the beats rapidly disappear with carbon dioxide and reappear under the influence of oxygen; and attention may be drawn to the fact that there is here no possible complication by action on nerve-fibres or on ganglion-cells, which do not yet exist. Submitted to tetanising currents, the embryonic, unlike the adult heart, is completely tetanised. Muscarine, which promptly abolishes the beat of an adult heart, has little or no action upon the embryonic heart.

According to Hermann and v. Gendre, the developing chick's embryo may exhibit an E.M.F. of  $\frac{1}{100}$  Dan., any part of the dorsal surface of the embryo being positive to any point of the yolk; thus, in the embryo itself the current is directed from ventral to dorsal surface.

Intra-uterine movements of the embryo may occur as early as the seventh week, although the obvious movements known to pregnant women as 'quickenings' are not usually noticed before the sixteenth week. Such movements are not voluntary but reflex, the result of internal changes in the composition of the blood or of external stimuli; according to Soltmann, the cortex cerebri of new-born puppies is inexcitable, and, as was mentioned on p. 495, the cortico-pyramidal system in man is of post-embryonic development. Anencephalous monsters, *i.e.* beings born without cerebrum or cerebellum, can perform reflex movements, and may even 'automatically' breathe, suck, cry, and swallow through the agency of bulbo-spinal centres.



## II. CONSTITUTIONAL FORMULÆ OF SOME OF THE CHIEF PROXIMATE PRINCIPLES

THE chemical relations of several of the more important substances mentioned in Chapters V and VI, such as the fats and many of the nitrogen compounds, will be better appreciated by considering their constitutional or linkage-formulæ, and the position that these formulæ occupy in well-known series. This point of view is particularly serviceable in the case of *urea*, *glycin*, *sarcosin*, *taurin*, *leucin*, and *tyrosin*, the constitutional formulæ of which can be given on a satisfactory basis; but it is not without value in the case of other complicated substances represented in a more or less hypothetical form, such as *uric acid*, *hippuric acid*, *creatin*, *creatinin*, *indican* (so-called), *indol*, *skatol*, *lecithin*, *cystin*; and the fats can be usefully so represented. As for the proteid group, and substances such as hæmoglobin, bile-pigments, &c., their graphic formulæ must be recognised as imaginary pictures containing radicles grouped so as to show possibilities, and arranged to suit a very scanty knowledge of facts.

The elements we have to deal with are hydrogen, oxygen, nitrogen, carbon, and occasionally phosphorus and sulphur.

The chief radicles entering into the composition of bodies that we have to consider are: hydroxyl,  $\text{HO}$ ; amidogen,  $\text{NH}_2$ ; the hydrocarbon radicles,  $\text{CH}_3$ ,  $\text{C}_2\text{H}_5$ ,  $\text{C}_3\text{H}_7$ ,  $\text{C}_3\text{H}_5$ , &c.; and the benzene radicles,  $\text{C}_6\text{H}_5$ ,  $\text{C}_6\text{H}_4$ , &c.

We shall best form an idea of their combinations by considering—

(1) Some simple linkage-formulæ of familiar bodies,  $\text{H}_2\text{O}$ ,  $\text{NH}_3$ ,  $\text{CO}_2$ ,  $\text{HNO}_3$ ,  $\text{H}_2\text{SO}_4$ ,  $\text{H}_3\text{PO}_4$ .

(2) The linkage-formulæ of a simple fatty acid series, of the combination of some of its members with the glycerin radicle,  $\text{C}_3\text{H}_5$ , as *fat*, and of two or three organic compounds with nitrogen, which from the chemical point of view fall into place on the lines of series running parallel with a simple fatty acid series.

(3) Linkage-formulæ that include a benzene nucleus,  $\text{C}_6\text{H}_5$ .

(4) Linkage-formulæ of a complicated or not definitely settled type.

The ordinary or empirical formulæ of water, ammonia, &c., translated into linked-formulæ, are as follows:

Water . . . . .  $\text{H}_2\text{O}$        $\text{H}-\text{O}-\text{H}$

Ammonia . . . . .  $\text{NH}_3$        $\begin{array}{c} \text{H}-\text{N}-\text{H} \\ | \\ \text{H} \end{array}$

Marsh gas . . . . .	CH <sub>4</sub>	$\begin{array}{c} \text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H} \end{array}$
Carbon dioxide . . . . .	CO <sub>2</sub>	O=C=O
Sulphuric acid . . . . .	H <sub>2</sub> SO <sub>4</sub>	$\begin{array}{c} \text{HO} \\   \\ \text{O}=\text{S}=\text{O} \\   \\ \text{HO} \end{array}$
Phosphoric acid . . . . .	H <sub>3</sub> PO <sub>4</sub>	$\begin{array}{c} \text{HO} \\   \\ \text{HO}-\text{P}-\text{HO} \\    \\ \text{O} \end{array}$

The formula of the hypothetical compound carbonic acid, H<sub>2</sub>CO<sub>3</sub>—which although not known as a free compound, but only as H<sub>2</sub>O + CO<sub>2</sub>, is indicated as a rational combination by the existence of corresponding salts, Na<sub>2</sub>CO<sub>3</sub>, K<sub>2</sub>CO<sub>3</sub>, &c.—serves as the starting-point of the series of fatty acids, and the type upon which urea is constructed. Its linked

formula is  $\text{HO}-\overset{\text{HO}}{\underset{|}{\text{C}}}=\text{O}$ ; that of urea is  $\text{NH}_2-\overset{\text{NH}_2}{\underset{|}{\text{C}}}=\text{O}$ ; that of acetone is  $\text{CH}_3-\overset{\text{CH}_3}{\underset{|}{\text{C}}}=\text{O}$ ; and substituting H for one of the hydroxyl radicles, HO, we

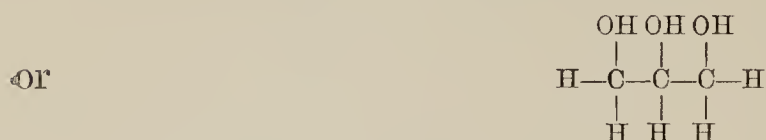
have  $\text{HO}-\overset{\text{H}}{\underset{|}{\text{C}}}=\text{O}$ , or formic acid, which is the starting-point of the fatty acid series, C<sub>n</sub>H<sub>2n+1</sub>. CO.OH, where CH<sub>2</sub>, or a multiple of CH<sub>2</sub>, is the difference between different members of the series.

Formic acid.	CH <sub>2</sub> O <sub>2</sub>	or	$\text{HO}-\overset{\text{H}}{\underset{ }{\text{C}}}=\text{O}$	or	H.CO.OH.
Acetic acid.	C <sub>2</sub> H <sub>4</sub> O <sub>2</sub>	or	$\begin{array}{c} \text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	or	CH <sub>3</sub> . CO.OH.
Propionic acid.	C <sub>3</sub> H <sub>6</sub> O <sub>2</sub>	or	$\begin{array}{c} \text{H} \\   \\ 2(\text{CH}_2) \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	or	C <sub>2</sub> H <sub>5</sub> . CO.OH.
Butyric acid.	C <sub>4</sub> H <sub>8</sub> O <sub>2</sub>	or	$\begin{array}{c} \text{H} \\   \\ 3(\text{CH}_2) \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	or	C <sub>3</sub> H <sub>7</sub> . CO.OH.
Caproic acid.	C <sub>6</sub> H <sub>12</sub> O <sub>2</sub>	or	$\begin{array}{c} \text{H} \\   \\ 5(\text{CH}_2) \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	or	C <sub>5</sub> H <sub>11</sub> . CO.OH.
Palmitic acid.	C <sub>16</sub> H <sub>32</sub> O <sub>2</sub>	or	$\begin{array}{c} \text{H} \\   \\ 15(\text{CH}_2) \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	or	C <sub>15</sub> H <sub>31</sub> . CO.OH.
Stearic acid.	C <sub>18</sub> H <sub>36</sub> O <sub>2</sub>	or	$\begin{array}{c} \text{H} \\   \\ 17(\text{CH}_2) \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	or	C <sub>17</sub> H <sub>35</sub> . CO.OH.

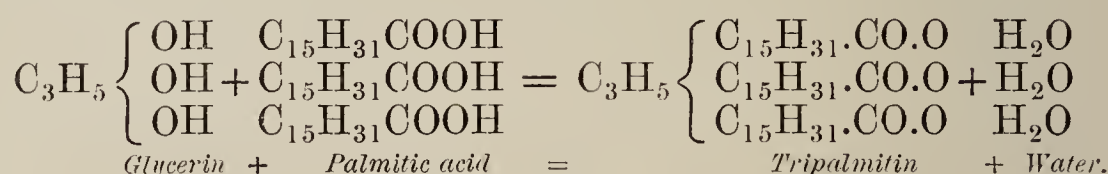


**Fatty acids, fats, and soaps.**—The animal fats are combinations of palmitic, stearic, and oleic acid with glycerin, in the form of tripalmitate, tristearate, and trioleate of glycerin.

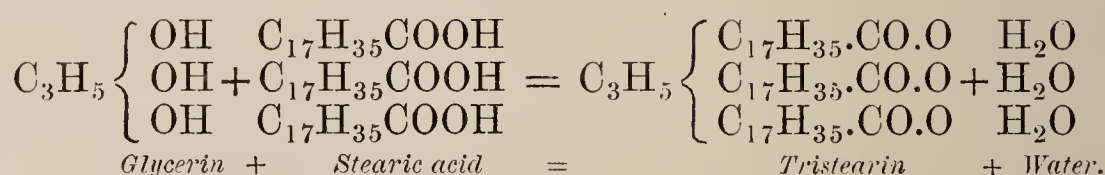
Glycerin is a triacid alcohol having the empirical formula  $C_3H_8O_3$ , the constitutional formula  $C_3H_5(OH)_3$ ,



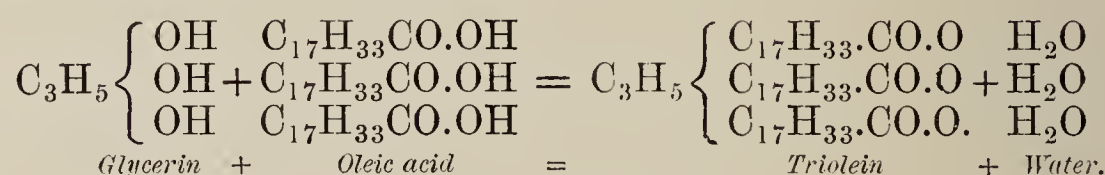
Palmitate of glycerin or *tripalmitin* is constituted as follows :



Stearate of glycerin or *tristearin* is constituted as follows :

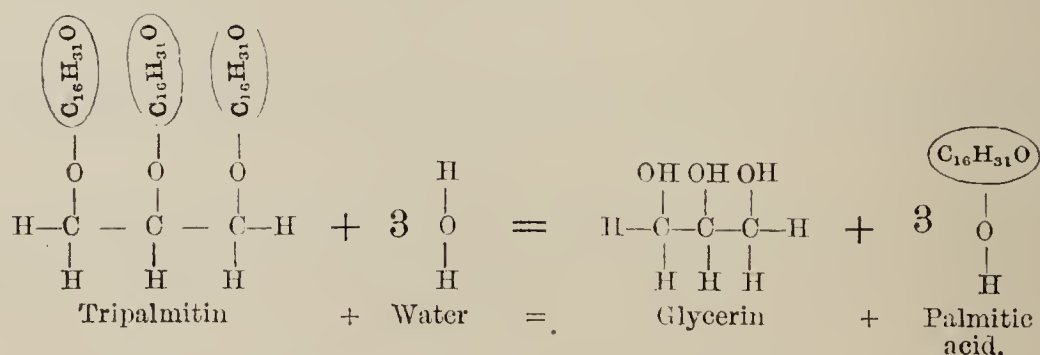


Oleate of glycerin or *triolein* is constituted as follows :



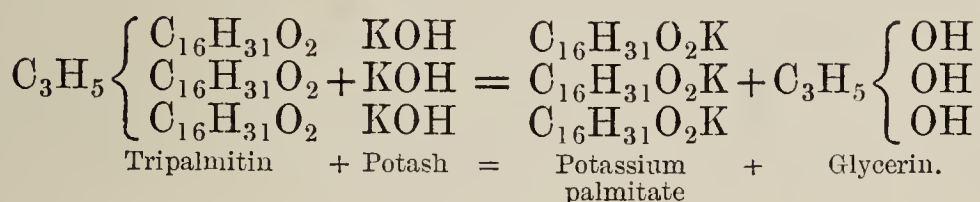
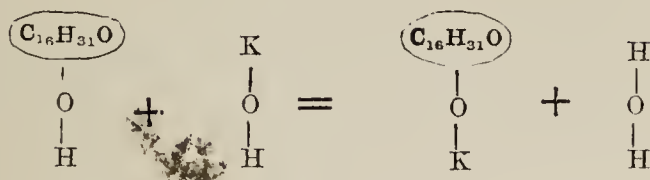
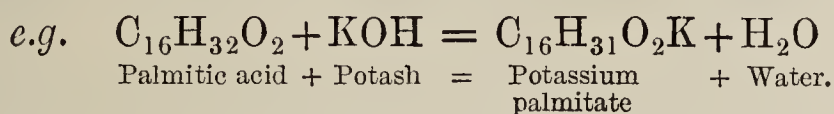
Animal fat is a mixture of the three glycerides—tripalmitin, tristearin, and triolein. Under the influence of ferments they undergo decomposition into glycerin and fatty acids.

These relations may be graphically represented as follows :



Fatty acids and fats are saponified by treatment with an alkali ;

soaps being compounds of the fatty acids with sodium or potassium, viz. palmitate, stearate, and oleate of sodium or potassium.



The analogous relations of the stearic and oleic compounds can be represented in a similar form, the palmitic radicle ( $C_{16}H_{31}O$ ) being replaced by the stearic radicle ( $C_{18}H_{35}O$ ) and the oleic radicle ( $C_{18}H_{33}O$ ).

**Glycin, sarcosin, lactic acid, leucin, tyrosin, taurin, and hippuric acid** are of parallel constitution with members of the fatty acid series already considered, as exhibited in the subjoined table, the amidogen radicle,  $NH_2$ , taking the place of H in the hydrocarbon radicle, or a hydrocarbon radicle replacing H in the amidogen radicle; tyrosin and hippuric acid include a benzoic nucleus, and belong therefore to aromatic bodies; in taurin, the sulphur group,  $SO_2.OH$ , replaces the carbon group,  $CO.OH$ .

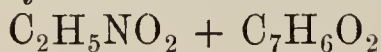
Formic acid.	Taurin or amido-ethyl-sulphonic acid.		
$\begin{array}{c} \text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{NH}_2 \\   \\ \text{H}-\text{C}-\text{H} \end{array}$	$\begin{array}{c} \text{H}-\text{C}-\text{H} \\   \\ \text{O}=\text{S}=\text{O} \\   \\ \text{OH} \end{array}$	
acid.	Glycin or amido-acetic acid.	Sarcosin or methyl-amido-acetic acid.	Hippuric acid or benzyl-amido-acetic acid.
$\begin{array}{c} \text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{NH}_2 \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{CH}_3 \\   \\ \text{N}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{C}_6\text{H}_5 \\   \\ \text{C}=\text{O} \\   \\ \text{N}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$



Propionic acid.	Sarcosine or $\alpha$ hydroxy-propionic acid.	Lactic acid or $\beta$ hydroxy-propionic acid	Tyrosin or hydroxy-phenyl-amido-propionic acid	Cystein or sulphamido-propionic acid
$\begin{array}{c} \text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{HO} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{HO} \\   \\ \text{C}_6\text{H}_4 \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{NH}_2 \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HS}-\text{C}-\text{NH}_2 \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$
Caproic acid.	Leucin or amido-caproic acid.			
$\begin{array}{c} \text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$	$\begin{array}{c} \text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{H} \\   \\ \text{H}-\text{C}-\text{NH}_2 \\   \\ \text{HO}-\text{C}=\text{O} \end{array}$			

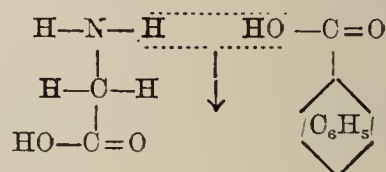
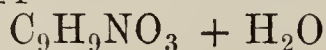
Other substances of physiological interest—in particular, uric acid and allied substances—being probably some of the chemical fragments of disintegrated proteid, cannot be arranged in definite series, and are in many cases of unknown or of doubtful structure. We shall, therefore, merely enumerate these bodies, giving their hypothetical constitutional formulæ, and drawing attention to the parallel relations of glycine and of taurine in hippuric and in the biliary acids.

Glycine and benzoic acid



form

Hippuric acid and water



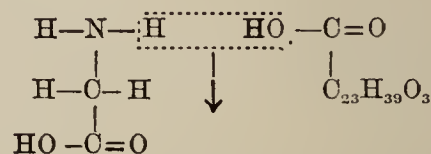
Similar conjugations are effected between glycine and cholic acid, and between taurine and cholic acid; but we have not the data for constructing a graphic formula of cholic acid: it probably contains, however, the group  $\text{HO}-\text{C}=\text{O}$ , and we may represent the relations as follows:

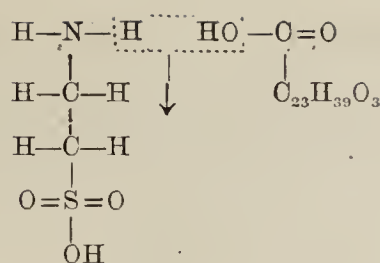
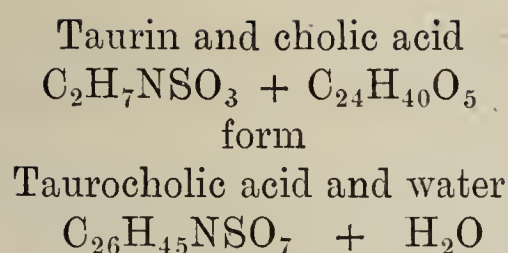
Glycine and cholic acid



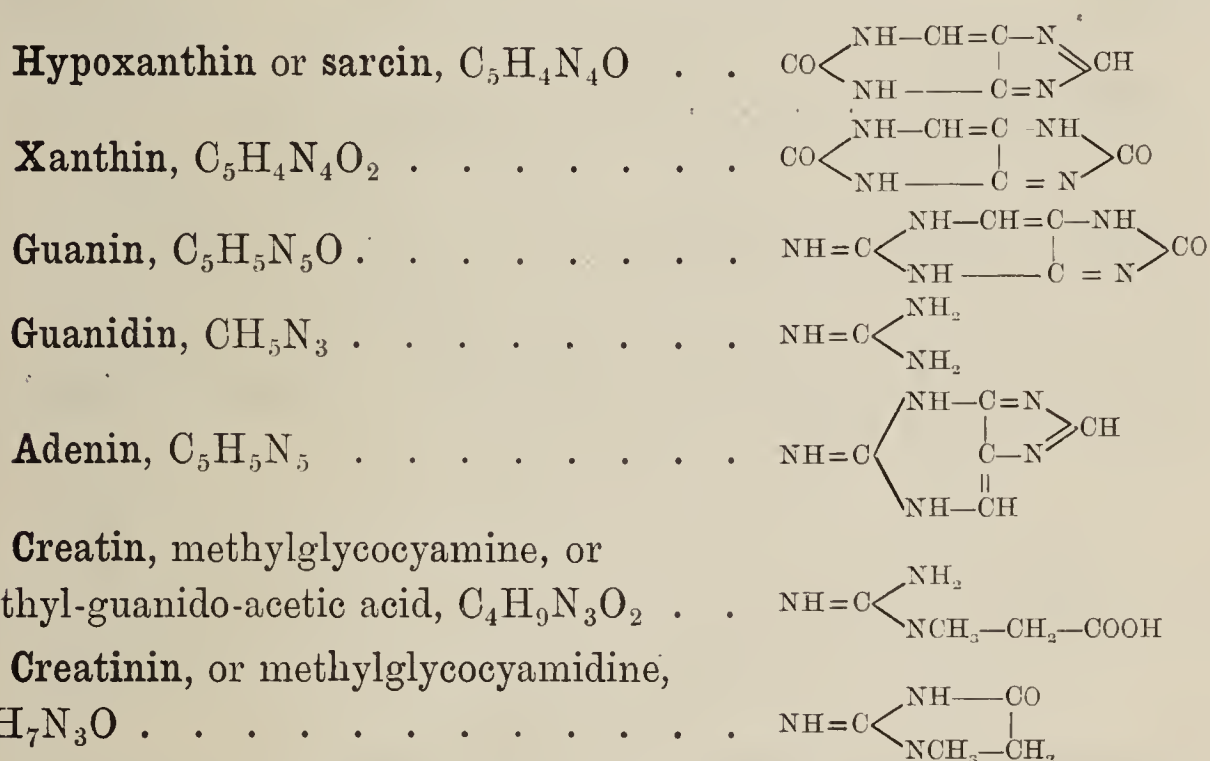
form

Glycocholic acid and water

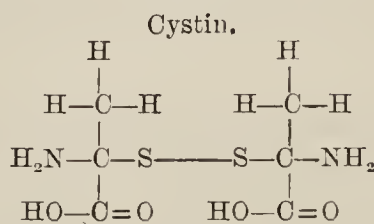
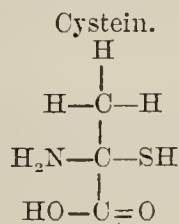
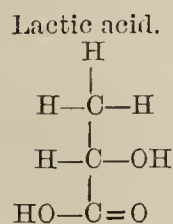




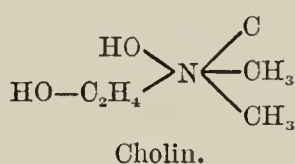
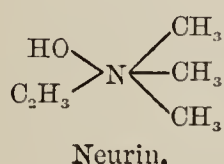
**Uric acid**,  $C_5H_4N_4O_3$ .—Of several schemata which have been proposed for uric acid, the most correct is probably that representing it as a diureide of acrylic acid:  $CO < \begin{array}{c} NH-CO-C-NH \\ NH \quad \quad \quad || \\ \quad \quad \quad C-NH \end{array} > CO$ . **Cynurenic acid**,  $C_{10}H_7NO_3 + H_2O$ ; **oxaluric acid**,  $C_3H_4N_2O_4$ ; **allantoin**,  $C_4H_6N_4O_3$ ; **xanthin**, **hypoxanthin**, **guanin**, and **adenin**, are bodies closely related to uric acid; the last four are related to nuclein, forming a group of nuclein bases.



**Cystein**,  $C_3H_7NSO_2$ , or sulphamido-lactic acid, and **cystin** ( $C_3H_6NSO_2$ )<sub>2</sub>, or disulphamido-lactic acid, are constituted on the lactic acid type; they are amido-lactic acids in which O is replaced by S.

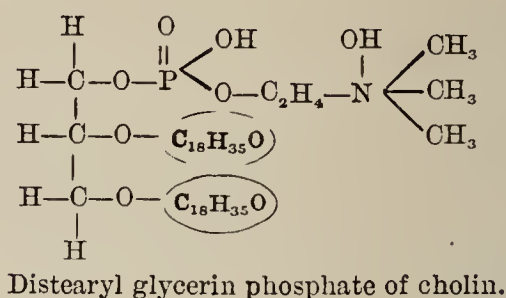
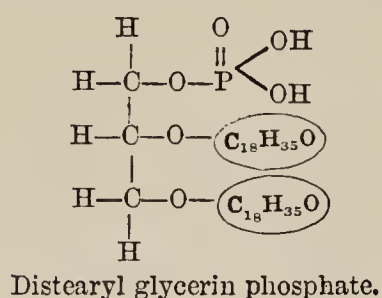


**Neurin**,  $C_5H_{13}NO$ , and **bilineurin** or **cholin**,  $C_5H_{15}NO_2$ , formerly regarded as being identical, are distinct bodies chemically allied to muscarin; neurin is poisonous, cholin is non-poisonous.



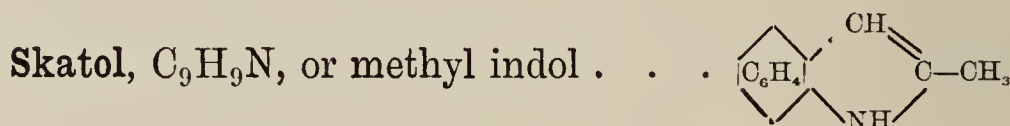


**Lecithin**,  $C_{44}H_{90}NPO_9$ . There are several lecithins or phosphorised fats; the most common is the distearyl glycerin phosphate of cholin, which may be represented as under:

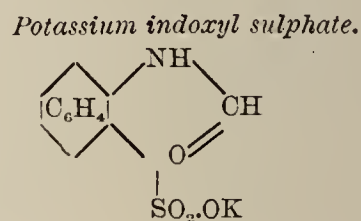
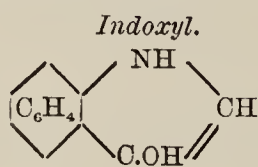


### *Aromatic Compounds*

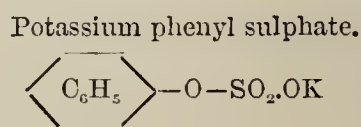
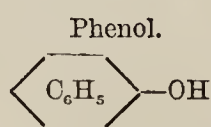
**Tyrosin** and **hippuric acid** have already been represented (pp. 593 and 594).



**Potassium indoxyl sulphate**,  $C_8H_6NKS O_4$ , so-called '*indican*' of urine, derived from indol:

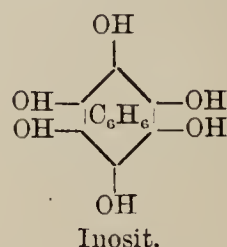
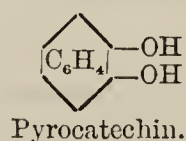


**Potassium phenyl sulphate**,  $C_6H_5KSO_4$ , derived from phenol:



These are the aromatic sulphates alluded to on p. 237.

**Pyrocatechin**,  $C_6H_6O_2$ , and **Inosit**,  $C_6H_{12}O_6$ , are not carbohydrates, but aromatic bodies. Pyrocatechin (orthodihydroxybenzol) is an oxidation-product of phenol, and reduces copper sulphate. Inosit (hexahydroxybenzol) is optically inactive, does not reduce copper sulphate, and undergoes lactic but not alcoholic fermentation.



*Carbohydrates*

The chief carbohydrates of physiological importance may be classified as follows (Tollens):

I. *Monosaccharids*,  $C_6H_{12}O_6$ —

1. Dextrose (glucose or grape sugar).    2. Lævulose (fruit sugar).    3. Galactose.

II. *Disaccharids*,  $C_{12}H_{22}O_{11}$ , splitting into two molecules of the Mono-group by hydrolytic fermentation—

1. Sucrose (cane sugar).    2. Lactose (milk sugar).    3. Maltose (malt sugar).  
 $+H_2O$      $+H_2O$      $+H_2O$   
 = Dextrose + Lævulose.    = Dextrose + Galactose.    = Dextrose + Dextrose.

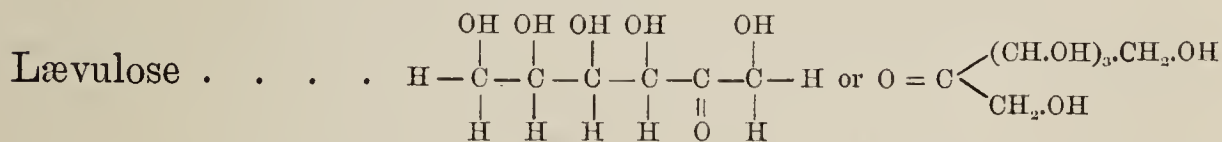
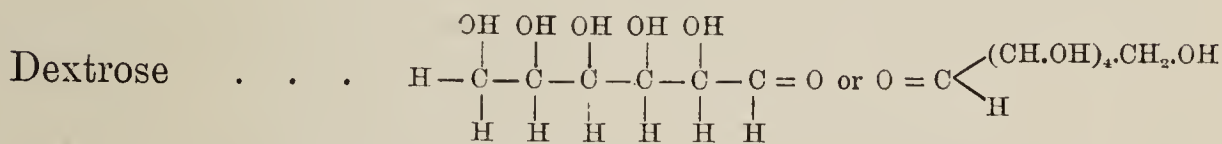
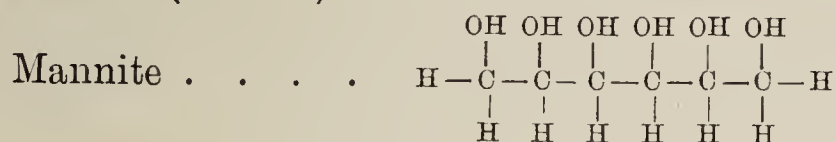
III. *Polysaccharids* ( $C_6H_{10}O_5$ )<sub>n</sub>, splitting into members of the Di- and Mono-groups by hydrolytic fermentation—

1. Starch.    2. Dextrin.    3. Glycogen.

The specific rotatory powers ( $\alpha_D$ ) of these several bodies are:

Dextrose	+52.7°	Sucrose	+64 to 66.5°	Starch	+197°
Lævulose	−80 to 85°	Lactose	+52.7°	Dextrin	+175 to 195°
Galactose	−20 to 25°	Maltose	+138°	Glycogen	+211 to 213°

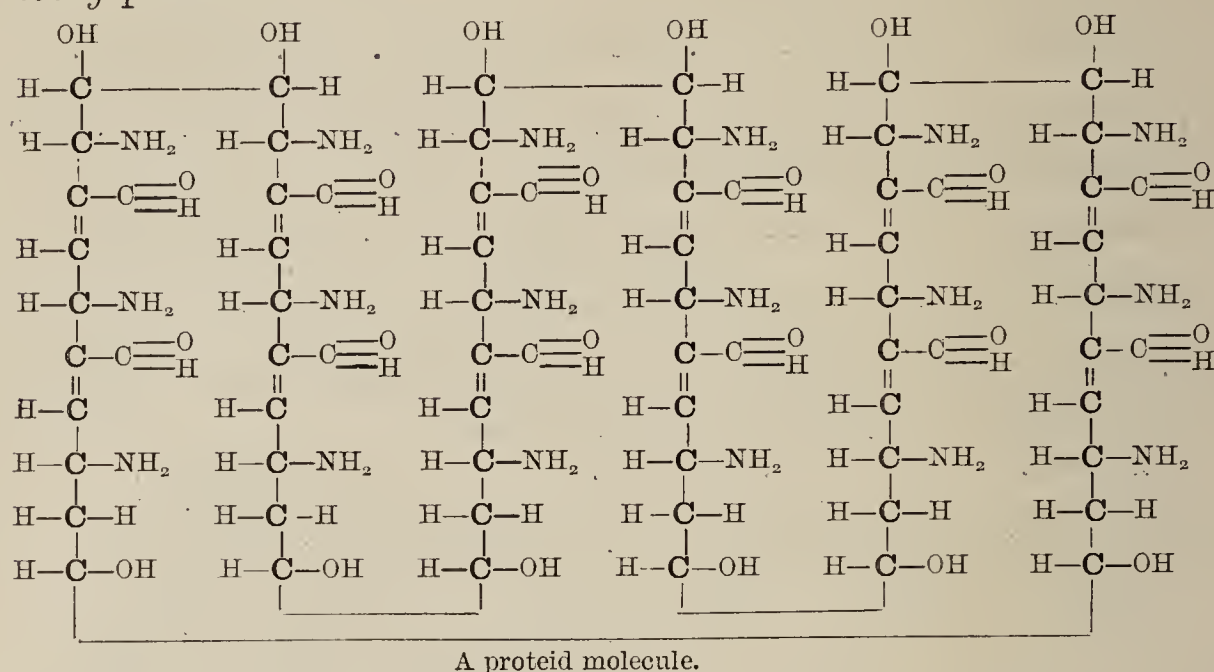
Dextrose and lævulose are closely related to the hex-acid alcohol, mannite,  $C_6H_8(OH)_6$ ; dextrose is constituted as an aldehyde, lævulose as a ketone (Fischer).



**Proteids.**—The structural formula of proteids is altogether conjectural, and that quoted below is given simply to emphasise the points alluded to in Chapter I, where we learned that proteid is the chemical centre of living matter—a large complex molecule which may take many shapes, and of which the constituent atoms or groups of atoms may ‘hold hands’ in many different ways, or hold out hands to other molecules with very different effects; a very slight shuffling of the molecule will convert a proteid food into a proteid poison, as we learned in the consideration of albumose (p. 188). Attention may be drawn to the conceivable combinations within the chemical range of such a molecule—(1) the amidogen group— $NH_2$ , in relation to urea and other



simple nitrogenous bodies ; (2) the excess of carbon and of hydrogen in relation to the deposition of carbohydrate or of fat ; (3) the *possibility* of a grouping into an aldehyde radicle,  $\text{H}-\text{C}=\text{O}$ , or into a cyanogen radicle  $-\text{CN}$ , in relation to the possible mode of action of *living* proteid.



### *Classification and Identification of Proteids*

Frequent allusion has been made to the various proteids found in the solids and fluids of the animal body, and the methods by which these proteids are separated and identified are described in their place under *Blood, Muscle, Urine, &c.*

The chief proteids met with as such in the animal body are the *albumins, globulins, and nucleo-albumins*. Several other bodies belonging to the proteid class do not exist ready formed in the living body, but are produced in the course of death changes—*fibrin* and *myosin* ; or in the process of digestion—*derived albumins, albumoses, and peptones* ; or by the manipulations used for their separation—*coagulated albumin*. In addition to these chief groups, there are numerous other bodies which, although resembling proteids in their chemical composition, are marked out by some distinguishing characteristic, and collectively spoken of under the title of *albuminoids* :—*mucin, elastin, chondrin, collagen, nuclein, gelatin*, and the pathological product *lardacein* belong to this group ; and *hæmoglobin*, although not classed with proteids, differs from them only by the presence of iron.

The classification of proteids is principally based upon their solubilities in distilled water, in dilute solutions of neutral salts, and in saturated saline solutions.

(1) *Albumins* are soluble in distilled water, in dilute and in saturated solutions of sodium or of magnesium sulphate, but insoluble in saturated solution of ammonium sulphate ; they can therefore be precipitated by saturating their solutions with the last-named salt.

(2) *Globulins* are insoluble in distilled water and in saturated saline solutions, but soluble in dilute saline solutions; they can therefore be precipitated from such solutions by removal of the salt by dialysis, or by saturation with a neutral salt, such as magnesium sulphate.

(3) *Nucleo-albumins* are very similar to globulins as regards solubility, but are chemically characterised by their high percentage of phosphorus; they are insoluble in distilled water, hardly soluble in dilute saline solutions, but soluble in dilute alkaline solution. They form part of cell-nuclei and cell-protoplasm ('cell-globulin'); on decomposition they yield *nuclein*, a phosphorised body originally obtained by the artificial digestion of pus cells.

(4) The digestion proteids—*acid-* and *alkali-albumin*, *albumoses*, and *peptones*—are characterised by their great solubility; *peptones* are soluble in all the media named above, and are not precipitated by saturation with ammonium sulphate; *albumoses* are soluble in dilute saline solutions, but insoluble in saturated ammonium sulphate solution; they are therefore precipitated from solution by saturation with this salt;<sup>1</sup> *acid-albumin* and *alkali-albumin* are soluble in dilute acid or alkaline solution, but insoluble in distilled water and in dilute saline; they are therefore precipitated by exact neutralisation of their solutions.

(5) The *post-mortem* proteids—*fibrin* and *myosin*—are comparatively insoluble; like the globulins, from which they are derived, they are insoluble in distilled water and in saturated saline solutions, but soluble in saline solutions of moderate strength—5 to 10 per cent.

(6) *Coagulated albumin* is insoluble in all these media; boiling with a strong acid, or warming with a weak acid, *plus* a digestive ferment, is required for its solution.

All proteids in solution give the following colour-reactions:—

1. *The xanthoproteic reaction*.—Boiling with nitric acid turns the solution to yellow, which is deepened to orange by the subsequent addition of strong ammonia or potash.

2. *Millon's reaction*.—Boiling with Millon's reagent (mercurous and mercuric nitrates) turns the solution red.

3. *The biuret reaction*.—A trace of copper sulphate and excess of caustic soda or potash give a violet or rose colour.

The first two tests applied to concentrated solutions give a precipitate turning yellow, orange, or red, as the case may be.

By means of heat-coagulation a proteid is known to belong to one or

<sup>1</sup> But it is now agreed that albumoses (deutero-albumose in particular) are not completely precipitated by ammonium sulphate. A solution must be raised to boiling-point and saturated, during alkaline, neutral and acid reaction, before the final filtrate may be assumed to contain only peptone and no albumose; this 'peptone' residue is a brownish hygroscopic substance, not giving any biuret reaction (Kühne). Pekelharing goes a step further, asserting that even this final 'peptone' is an albumose mixture. The family name, 'peptones,' formerly borne by the digestion proteids, has gradually given way to the family name, 'albumoses.'



other of the following groups—(a) to the albumins or globulins, which coagulate, or (b) to the derived albumins, albumoses, or peptones, which are not coagulated by heat.

The proteids coagulable by heat are distinguished into albumins and globulins by saturation with magnesium sulphate; globulins are, albumins are not, precipitated, and if both are present, globulins can be retained on a filter-paper, while albumins pass through in the filtrate.

The two chief forms of albumin ordinarily met with are egg-albumin and serum-albumin; the first of these is, the second is not, coagulated by shaking with ether. Muscle-albumin is practically indistinguishable from serum-albumin. The globulin class includes serum-globulin and fibrinogen, and is widely distributed in the solid organs, *e.g.* in muscle as muscle-globulin, in the crystalline lens as crystallin. We do not in ordinary analysis attempt to distinguish between different members of this class, although actually they have been distinguished by varying solubilities in saline solutions, and by their behaviour with fibrin-ferment. The *post-mortem* proteids—fibrin and myosin—are also included in the globulin class. They are to some extent distinguished by their solubilities in saline solutions, and by the temperature at which they coagulate.

To further distinguish between and identify the group of proteids that are not coagulable by heat, we must examine—(1) the reaction, (2) the character of the biuret colour, (3) the behaviour to saturation with ammonium sulphate.

Acid-albumin in solution has an acid reaction, and is precipitated by cautious neutralisation, excess of alkali being avoided.

Alkali-albumin in solution has an alkaline reaction, and is likewise precipitated by cautious neutralisation with acid.

Albumoses are precipitated by saturation with ammonium sulphate.<sup>1</sup> peptones are not. If a proteid fluid that is not coagulable by heat, is neutralised exactly and thrown on a filter, acid- or alkali-albumins are retained, albumoses and peptones pass in the filtrate; if this filtrate is saturated with ammonium sulphate and filtered, albumoses are retained and peptones pass through; the presence of peptones is recognised in this last filtrate by a rose-coloured biuret reaction, care being taken to add KHO in excess.

*Summary of the Principal Proteids and of their Distinguishing Tests*

Coag. on boiling	{	Egg-albumin . . .	Ether ppt.
		Serum-albumin . . .	No ether ppt. No MgSO <sub>4</sub> ppt.
		Globulins . . .	MgSO <sub>4</sub> ppt.
No coag. on boiling	{	Acid and alkali albumins	Neutralisation ppt.
		Albumoses . . .	HNO <sub>3</sub> ppt. in cold, sol. by heat.
		Peptones . . .	Pink biuret reaction in filtrate after saturation with Am <sub>2</sub> SO <sub>4</sub> .

<sup>1</sup> Cf. footnote on p. 599.

*Summary of the Digestion Proteids*

Test	Proto-Albumose	Hetero-Albumose	Deutero-Albumose	Peptone
<i>Biuret</i> ( $\text{CuSO}_4, \text{KHO}$ )	Pink	Pink	Pink	Pink
<i>Water, hot and cold</i> .	Soluble	Insoluble ; ppt. by di- alysis	Soluble	Soluble
<i>Saturation by NaCl or MgSO<sub>4</sub></i>	ppt.	ppt.	no ppt. (ppt. by dilute HA)	no ppt.
<i>Saturation by Am<sub>2</sub>SO<sub>4</sub></i>	ppt.	ppt.	ppt. (incompl.)	no ppt.
<i>Nitric Acid</i> . . .	ppt. in cold diss. by heat returns in cold	Do.	Do. in pre- sence of excess NaCl	no ppt.
<i>CuSO<sub>4</sub></i> . . .	ppt.	ppt.	no ppt.	no ppt.
<i>Diffusibility</i> . .	moderate	small	moderate	great

A gastric digest of fibrin, made as described on pp. 187 and 188 (after removal of heat coagulum and neutralisation ppt., if any), contains the three albumoses and peptones; in addition to the pink biuret reaction, it should give the characteristic nitric acid reaction, and on dropping in distilled water a cloud should form (hetero-albumose).

They may be roughly separated as follows (Halliburton) :—

*Precipitate by saturation with Am<sub>2</sub>SO<sub>4</sub> and filter.*

Albumoses on filter.....Peptones in filtrate.

*Dissolve in water, precipitate by saturation with NaCl, and filter.*

Proto- and hetero-albumose on filter...Deutero-albumose in filtrate.

*Dissolve in water, dialyse, and filter. Ppt. by alcohol, wash and dry.*

Hetero-albumose on filter.....Proto-albumose in filtrate.

*Wash and dry.*

*Ppt. by alcohol, wash and dry.*



## III. UNITS OF MEASUREMENT

## FRENCH MEASURES OF LENGTH, WEIGHT, AND VOLUME

$$1 \text{ millimètre} = \frac{1}{10} \text{ centimètre} = \frac{1}{1000} \text{ mètre.}$$

$$1 \text{ centimètre} = \frac{1}{100} \text{ mètre.}$$

$$1 \text{ micromillimètre (1 } \mu) = \frac{1}{1000} \text{ millimètre.}$$

$$1 \text{ litre} = 1000 \text{ centimètres cubes (1000 c.c.).}$$

*Relation between Volume and Weight*

$$1 \text{ c.c.} = 1 \text{ gramme.}$$

$$1 \text{ milligramme} = \frac{1}{1000} \text{ gramme.}$$

$$1 \text{ kilogramme} = 1000 \text{ grammes.}$$

## RELATIONS BETWEEN ENGLISH AND FRENCH MEASURES

*Length. English to French*

$$1 \text{ inch} = 2.539954 \text{ centimètres.}$$

$$1 \text{ foot} = 30.479449 \text{ ,,}$$

$$1 \text{ yard} = 91.438347 \text{ ,,}$$

$$1 \text{ mile} = 1609.315 \text{ mètres}$$

(To convert inches into centimètres multiply by  $\frac{25}{10}$ .)

*Length. French to English*

$$1 \text{ mètre} = 39.37079 \text{ inches.}$$

$$1 \text{ centimètre} = 0.39371 \text{ inch.}$$

$$1 \text{ micromillimètre} = 0.00003937 \text{ inch.}$$

(To convert centimètres into inches multiply by  $\frac{10}{25}$ .)

*Weight. English to French*

$$1 \text{ grain} = 0.0648 \text{ gramme.}$$

$$1 \text{ ounce} = 28.3495 \text{ grammes.}$$

$$1 \text{ pound} = 453.592 \text{ ,,}$$

$$1 \text{ stone} = 6.35 \text{ kilogrammes.}$$

$$1 \text{ cwt.} = 50.8 \text{ ,,}$$

$$1 \text{ ton} = 1016 \text{ ,,}$$

*Weight. French to English*

$$1 \text{ gramme} = 15.432349 \text{ grains.}$$

$$1 \text{ kilogramme} = 2.2046213 \text{ pounds,}$$

or about 35 ounces.

$$1 \text{ milligramme} = 0.015432 \text{ grain.}$$

*Volume. English to French*

$$1 \text{ cubic inch} = 16.3861759 \text{ centimètres cubes.}$$

$$1 \text{ fluid ounce} = 28.3495 \text{ ,,}$$

$$1 \text{ pint} = 567 \text{ ,,}$$

$$1 \text{ cubic foot} = 28.3153 \text{ litres.}$$

*Volume. French to English*

1 centimètre cube	=	0·061027 cubic inch.
1 litre (1000 c.c.)	=	61·027 cubic inches, or 35 fluid ounces, or $1\frac{3}{4}$ pints.
1 mètre cube (1000 litres)	=	35·3 cubic feet.

*Measures of Surface*

1 square mètre	=	about 1550 square inches, or 10,000 square centimètres, or 10·75 square feet.
1 square inch	=	about 6·45 square centimètres.
1 square foot	=	about 930 „ „

*Measures of Energy*

1 kilogrammètre	=	about 7·24 foot-pounds = 98,100,000 ergs.
1 foot-pound	=	„ 0·1381 KgM.
1 foot-ton	=	„ 310 KgMs.
1 milligrammètre	=	98·1 ergs.

*Mechanical Equivalent of Heat*

1 kilocalorie	=	424 (or 423·985) kilogrammètres = 41,593,010,000 ergs.
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*Approximate Solubilities in Water at 15° to 18° C. of some Crystallised Salts in Frequent Use*

Sodium chloride	.	.	.	.	36 per 100
Sodium sulphate	.	.	.	.	50 „
Ammonium chloride	.	.	.	.	36 „
Ammonium sulphate	.	.	.	.	50 „
Magnesium sulphate	.	.	.	.	125 „
Zinc sulphate	.	.	.	.	160 „

<i>Volumes of</i>	<i>Weights of</i>
1 gram. hydrogen = 11160 c.c.	1000 c.c. hydrogen = 0·0895 gram.
„ oxygen = 697 „	„ oxygen = 1·43 „
„ nitrogen = 797 „	„ nitrogen = 1·25 „
„ carbon di-	„ carbon di-
oxide = 507 „	oxide = 1·97 „
„ air = 775 „	„ air = 1·29 „
„ water-	„ water-vapour = 0·82 „
vapour = 1240 „	

*Coefficient of expansion of gases,  $\frac{1}{273}$ , or 0·00366 per 1° C.*

*Area of circle* =  $\pi r^2$ , or  $3·1416r^2$ .

*Surface of sphere* =  $4\pi r^2$ , or  $12·5664r^2$ .

*Volume of sphere* =  $\frac{4\pi r^3}{3}$ , or  $4·2r^3$ .



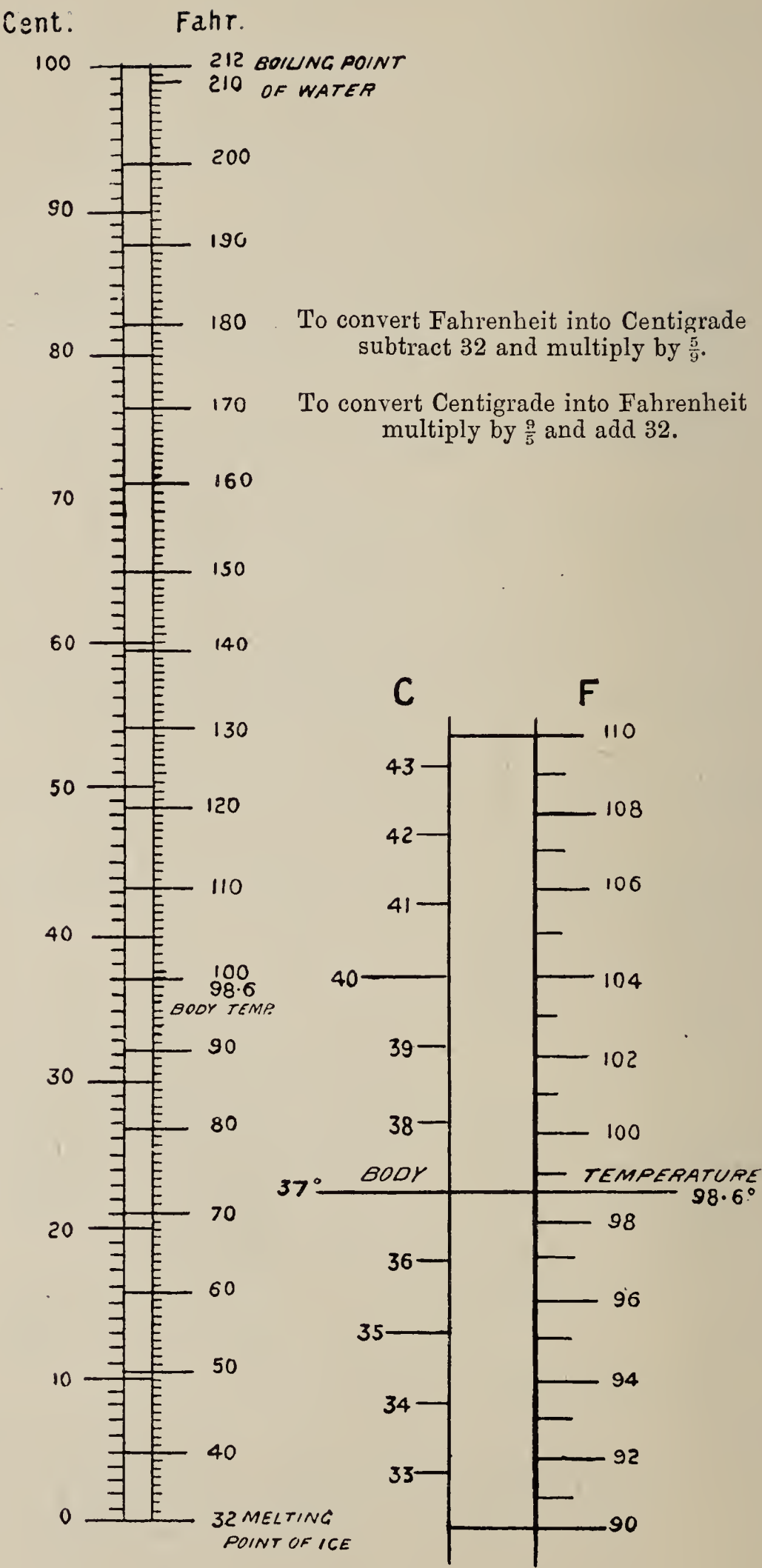


FIG. 315.—CENTIGRADE AND FAHRENHEIT THERMOMETER SCALES.

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